

UTILITY  
PATENT APPLICATION  
TRANSMITTAL

(Only for new nonprovisional applications under 37 CFR 1.53(b))

Attorney Docket No.

4085-226-27

First Inventor or Application Identifier

Michelle A.J. PALMER, et al

Title

RECEPTOR FUNCTION ASSAY FOR G-PROTEIN COUPLED RECEPTORS  
AND ORPHAN RECEPTORS BY REPORTER ENZYME MUTANT  
COMPLEMENTATION

APPLICATION ELEMENTS

See MPEP chapter 600 concerning utility patent application contents

ADDRESS TO:

Assistant Commissioner for Patents  
Box Patent Application  
Washington, DC 20231

1. ☒ Fee Transmittal Form (e.g. PTO/SB/17)  
(Submit an original and a duplicate for fee processing)
2. ☒ Specification Total Pages **25**
3. ☒ Drawing(s) (35 U.S.C. 113) Total Sheets **66**
4. ☐ Oath or Declaration Total Pages 
  - a. ☐ Newly executed (original or copy)
  - b. ☐ Copy from a prior application (37 C.F.R. §1.63(d))  
(for continuation/divisional with box 15 completed)
    - i. ☐ DELETION OF INVENTOR(S)  
Signed statement attached deleting inventor(s) named  
in the prior application, see 37 C.F.R. §1.63(d)(2) and  
1.33(b).
5. ☐ Incorporation By Reference (usable if box 4B is checked)  
The entire disclosure of the prior application, from which a copy of  
the oath or declaration is supplied under Box 4B, is considered to be  
part of the disclosure of the accompanying application and is hereby  
incorporated by reference therein.

ACCOMPANYING APPLICATION PARTS

6. ☐ Assignment Papers (cover sheet & document(s))
7. ☐ 37 C.F.R. §3.73(b) Statement ☐ Power of Attorney  
(when there is an assignee)
8. ☐ English Translation Document (if applicable)
9. ☐ Information Disclosure Statement (IDS)/PTO-1449 ☐ Copies of IDS Citations
10. ☐ Preliminary Amendment
11. ☒ White Advance Serial No. Postcard
12. ☐ Small Entity Statement(s) ☐ Statement filed in prior application. Status still proper and desired.
13. ☐ Certified Copy of Priority Document(s)  
(if foreign priority is claimed)
14. ☒ Other: List of Inventors' Names and Addresses

15. If a CONTINUING APPLICATION, check appropriate box, and supply the requisite information below:

☐ Continuation ☐ Divisional ☐ Continuation-in-part (CIP) of prior application no.:  
Prior application information: Examiner: Group Art Unit:

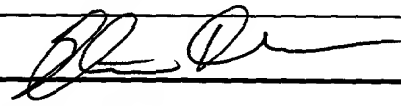
16. Amend the specification by inserting before the first line the sentence:

☐ This application is a ☐ Continuation ☐ Division ☐ Continuation-in-part (CIP)  
of application Serial No. Filed on

☒ This application claims priority of provisional application Serial No. 60/180,669 Filed February 7, 2000

17. CORRESPONDENCE ADDRESS

Steven B. Kelber  
PIPER MARBURY RUDNICK & WOLFE LLP  
1200 Nineteenth Street, N.W.  
Washington, D.C. 20036-2412  
Telephone No. (202) 861-3900  
Facsimile No. (202) 223-2085

Name:	Steven B. Kelber	Registration No.:	30,073
Signature:		Date:	9/11/00
Name:	Laura A. Donnelly	Registration No.:	38,435

Docket No. 4085-226-27

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

INVENTOR(S) Michelle A.J. PALMER, et al

SERIAL NO: New Application

FILING DATE: Herewith

FOR: RECEPTOR FUNCTION ASSAY FOR G-PROTEIN COUPLED RECEPTORS AND ORPHAN RECEPTORS BY REPORTER ENZYME MUTANT COMPLEMENTATION

FEE TRANSMITTAL

ASSISTANT COMMISSIONER FOR PATENTS  
WASHINGTON, D.C. 20231

FOR	NUMBER FILED	NUMBER EXTRA	RATE	CALCULATIONS
TOTAL CLAIMS	30 - 20 =	10	× \$18 =	\$180.00
INDEPENDENT CLAIMS	8 - 3 =	5	× \$78 =	\$390.00
<input type="checkbox"/> MULTIPLE DEPENDENT CLAIMS (If applicable)			+ \$260 =	\$0.00
<input checked="" type="checkbox"/> LATE FILING OF DECLARATION			+ \$130 =	\$130.00
BASIC FEE				\$690.00
TOTAL OF ABOVE CALCULATIONS				\$1,390.00
<input type="checkbox"/> REDUCTION BY 50% FOR FILING BY SMALL ENTITY				\$0.00
<input type="checkbox"/> FILING IN NON-ENGLISH LANGUAGE			+ \$130 =	\$0.00
<input type="checkbox"/> RECORDATION OF ASSIGNMENT			+ \$40 =	\$0.00
TOTAL				\$1,390.00

- ☐ Please charge Deposit Account No. 50-1442 in the amount of . A duplicate copy of this sheet is enclosed.
- ☒ A check in the amount of \$1,390.00 to cover the filing fee is enclosed.
- ☐ The Commissioner is hereby authorized to charge any additional fees which may be required for the papers being filed herewith and for which no check is enclosed herewith, or credit any overpayment to Deposit Account No. 50-1442. A duplicate copy of this sheet is enclosed.

Respectfully submitted,

PIPER MARBURY RUDNICK & WOLFE LLP



Steven B. Kelber  
Registration No. 30,073

Laura A. Donnelly  
Registration No. 38,435

Date:

9/1/02

1200 Nineteenth Street, N.W.  
Washington, D.C. 20036-2412  
Telephone No. (202) 861-3900  
Facsimile No. (202) 223-2085

**TITLE OF THE INVENTION**

**RECEPTOR FUNCTION ASSAY FOR G-PROTEIN COUPLED RECEPTORS AND ORPHAN RECEPTORS BY REPORTER ENZYME MUTANT COMPLEMENTATION**

5

**BACKGROUND OF THE INVENTION**

This application claims the benefit from Provisional Application Serial No. 60/180,669, filed February 7, 2000. The entirety of that provisional application is incorporated herein by reference.

10

**Field of the Invention**

This invention relates to methods of detecting G-protein-coupled receptor (GPCR) activity, and provides methods of assaying GPCR activity and methods for screening for GPCR ligands, G-protein-coupled receptor kinase (GRK) activity, and compounds that interact with components of the GPCR regulatory process.

15

The actions of many extracellular signals are mediated by the interaction of G-protein-coupled receptors (GPCRs) and guanine nucleotide-binding regulatory proteins (G-proteins). G-protein-mediated signaling systems have been identified in many divergent organisms, such as mammals and yeast. The GPCRs represent a large super family of proteins which have divergent amino acid sequences, but share common structural features, in particular, the presence of seven transmembrane helical domains. GPCRs respond to, among other extracellular signals, neurotransmitters, hormones, odorants and light. Individual GPCR types activate a particular signal transduction pathway; at least ten different signal transduction pathways are known to be activated via GPCRs. For example, the beta 2-adrenergic receptor ( $\beta 2AR$ ) is a prototype mammalian GPCR. In response to agonist binding,  $\beta 2AR$  receptors activate a G-protein (Gs) which in turn stimulates adenylate cyclase activity and results in increased cyclic adenosine monophosphate (cAMP) production in the cell.

20

25

The signaling pathway and final cellular response that result from GPCR stimulation depends on the specific class of G-protein with which the particular receptor is coupled (Hamm, "The many faces of G-Protein Signaling." J. Biol. Chem., 273:669-672 (1998)). For instance, coupling to the Gs class of G-proteins stimulates cAMP production and activation of Protein Kinase A and C pathways, whereas coupling to the Gi class of G-proteins down regulates cAMP. Other second messenger systems as calcium, phospholipase C, and phosphatidylinositol 3 may also be utilized. As a consequence, GPCR signaling events have predominantly been measured via quantification of these second messenger products.

A common feature of GPCR physiology is desensitization and recycling of the receptor through the processes of receptor phosphorylation, endocytosis and dephosphorylation (Ferguson, et al., "G-protein-coupled receptor regulation: role of G-protein-coupled receptor kinases and arrestins." Can. J. Physiol. Pharmacol., 74:1095-1110 (1996)). Ligand-occupied GPCRs can be phosphorylated by two families of serine/threonine kinases, the G-protein-coupled receptor kinases (GRKs) and the second messenger-dependent protein kinases such as protein kinase A and protein kinase C. Phosphorylation by either class of kinases serves to down-regulate the receptor by uncoupling it from its corresponding G-protein. GRK-phosphorylation also serves to down-regulate the receptor by recruitment of a class of proteins known as the arrestins that bind the cytoplasmic domain of the receptor and promote clustering of the receptor into endocytic vesicles. Once the receptor is endocytosed, it will either be degraded in lysosomes or dephosphorylated and recycled back to the plasma membrane as fully-functional receptor.

Binding of an arrestin protein to an activated receptor has been documented as a common phenomenon for a variety of GPCRs ranging from rhodopsin to  $\beta$ 2AR to the

neurotensin receptor (Barak, et al., "A  $\beta$ -arrestin/Green Fluorescent fusion protein biosensor for detecting G-Protein-Coupled Receptor Activation," J. Biol. Chem., 272:27497-500 (1997)). Consequently, monitoring arrestin interaction with a specific GPCR can be utilized as a generic tool for measuring GPCR activation. Similarly, a single G-protein and GRK also partner with a variety of receptors (Hamm, et al. (1998) and Pitcher et al., "G-Protein-Coupled Receptor Kinases," Annu. Rev. Biochem., 67:653-92 (1998)), such that these protein/protein interactions may also be monitored to determine receptor activity.

The present invention involves the use of a proprietary technology (ICAST<sup>TM</sup>, Intercistronic Complementation Analysis Screening Technology) for monitoring protein/protein interactions in GPCR signaling. The method involves using two inactive  $\beta$ -galactosidase mutants, each of which is fused with one of two interacting protein pairs, such as a GPCR and an arrestin. The formation of an active  $\beta$ -galactosidase complex is driven by interaction of the target proteins. In this system,  $\beta$ -galactosidase activity acts as a read out of GPCR activity. FIGURE 23 is a schematic depicting the method of the present invention. FIGURE 23 shows two inactive mutants that become active when they interact. In addition, this technology could be used to monitor GPCR-mediated signaling pathways via other downstream signaling components such as G-proteins, GRKs or c-Src.

Many therapeutic drugs in use today target GPCRs, as they regulate vital physiological responses, including vasodilation, heart rate, bronchodilation, endocrine secretion and gut peristalsis. See, e.g., Lefkowitz et al., Annu. Rev. Biochem., 52:159 (1983). For instance, drugs targeting the highly studied GPCR,  $\beta$ 2AR are used in the treatment of anaphylaxis, shock hypertension, asthma and other conditions. Some of these drugs mimic

the ligand for this receptor. Other drugs act to antagonize the receptor in cases when disease arises from spontaneous activity of the receptor.

Efforts such as the Human Genome Project are identifying new GPCRs ("orphan" receptors) whose physiological roles and ligands are unknown. It is estimated that several thousand GPCRs exist in the human genome. Of the 250 GPCRs identified to date, only 150 have been associated with ligands.

### **SUMMARY OF THE INVENTION**

A first aspect of the present invention is a method that monitors GPCR function proximally at the site of receptor activation, thus providing more information for drug discovery purposes due to fewer competing mechanisms. Activation of the GPCR is measured by a read-out for interaction of the receptor with a regulatory component such as arrestin, G-protein, GRK or other kinases, the binding of which to the receptor is dependent upon agonist occupation of the receptor. Protein/protein interaction is detected by complementation of reporter proteins such as utilized by the ICAST™ technology.

A further aspect of the present invention is a method of assessing G-protein-coupled receptor (GPCR) pathway activity under test conditions by providing a test cell that expresses a GPCR, e.g., muscarinic, adrenergic, dopamine, angiotensin or endothelin, as a fusion protein to a mutant reporter protein and interacting, i.e., G-proteins, arrestin or GRK, as a fusion protein with a complementing reporter protein. When test cells are exposed to a known agonist to the target GPCR under test conditions, activation of the GPCR will be monitored by complementation of the reporter enzyme. Increased reporter enzyme activity reflects interaction of the GPCR with its interacting protein partner.

A further aspect of the present invention is a method of assessing GPCR pathway activity in the presence of a test kinase.

A further aspect of the present invention is a method of assessing GPCR pathway activity in the presence of a test G-protein.

5 A further aspect of the present invention is a method of assessing GPCR pathway activity upon exposure of the test cell to a test ligand.

A further aspect of the present invention is a method of assessing GPCR pathway activity upon co-expression in the test cell of a second receptor.

10 A further aspect of the present invention is a method for screening for a ligand or agonists to an orphan GPCR. The ligand or agonist could be contained in natural or synthetic libraries or mixtures or could be a physical stimulus. A test cell is provided that expresses the orphan GPCR as a fusion protein with one  $\beta$ -galactosidase mutant and, for example, an arrestin or mutant form of arrestin as a fusion protein with another  $\beta$ -galactosidase mutant. The interaction of the arrestin with the orphan GPCR upon receptor activation is measured by  
15 enzymatic activity of the complemented  $\beta$ -galactosidase. The test cell is exposed to a test compound, and an increase in  $\beta$ -galactosidase activity indicates the presence of a ligand or agonist.

A further aspect of the present invention is a method for screening a protein of interest, for example, an arrestin protein (or mutant form of the arrestin protein) for the ability  
20 to bind to a phosphorylated, or activated, GPCR. A cell is provided that expresses a GPCR and contains  $\beta$ -arrestin. The cell is exposed to a known GPCR agonist and then reporter enzyme activity is detected. Increased reporter enzyme activity indicates that the  $\beta$ -arrestin molecule can bind to phosphorylated, or activated, GPCR in the test cell.

A further aspect of the present invention is a method to screen for an agonist to a specific GPCR. The agonist could be contained in natural or synthetic libraries or could be a physical stimulus. A test cell is provided that expresses a GPCR as a fusion protein with one  $\beta$ -galactosidase mutant and, for example, an arrestin as a fusion protein with another  $\beta$ -galactosidase mutant. The interaction of arrestin with the GPCR upon receptor activation is measured by enzymatic activity of the complemented  $\beta$ -galactosidase. The test cell is exposed to a test compound, and an increase in  $\beta$ -galactosidase activity indicates the presence of an agonist. The test cell may express a known GPCR or a variety of known GPCRs, or may express an unknown GPCR or a variety of unknown GPCRs. The GPCR may be, for example, an odorant GPCR or a  $\beta$ AR GPCR.

A further aspect of the present invention is a method of screening a test compound for G-protein-coupled receptor (GPCR) antagonist activity. A test cell is provided that expresses a GPCR as a fusion protein with one  $\beta$ -galactosidase mutant and, for example, an arrestin as a fusion protein with another  $\beta$ -galactosidase mutant. The interaction of arrestin with the GPCR upon receptor activation is measured by enzymatic activity of the complemented  $\beta$ -galactosidase. The test cell is exposed to a test compound, and an increase in  $\beta$ -galactosidase activity indicates the presence of an agonist. The cell is exposed to a test compound and to a GPCR agonist, and reporter enzyme activity is detected. When exposure to the agonist occurs at the same time as or subsequent to exposure to the test compound, a decrease in  $\beta$ -galactosidase activity after exposure to the test compound indicates that the test compound has antagonist activity to the GPCR.

A further aspect of the present invention is a method of screening a sample solution for the presence of an agonist, antagonist or ligand to a G-protein-coupled receptor (GPCR).



A test cell is provided that expresses a GPCR fusion and contains, for example, a  $\beta$ -arrestin protein fusion. The test cell is exposed to a sample solution, and reporter enzyme activity is assessed. Changed reporter enzyme activity after exposure to the sample solution indicates the sample solution contains an agonist, antagonist or ligand for a GPCR expressed in the cell.

5 A further aspect of the present invention is a method of screening a cell for the presence of a G-protein-coupled receptor (GPCR).

A further aspect of the present invention is a method of screening a plurality of cells for those cells which contain a G-protein coupled receptor (GPCR).

10 A further aspect of the invention is a method for mapping GPCR-mediated signaling pathways. For instance, the system could be utilized to monitor interaction of c-src with  $\beta$ -arrestin-1 upon GPCR activation. Additionally, the system could be used to monitor protein/protein interactions involved in cross-talk between GPCR signaling pathways and other pathways such as that of the receptor tyrosine kinases or Ras/Raf.

15 A further aspect of the invention is a method for monitoring homo- or hetero-dimerization of GPCRs upon agonist or antagonist stimulation.

20 A further aspect of the invention is a method of screening a cell for the presence of a G-protein-coupled receptor (GPCR) responsive to a GPCR agonist. A cell is provided that contains protein partners that interact downstream in the GPCR's pathway. The protein partners are expressed as fusion proteins to the mutant, complementing enzyme and are used to monitor activation of the GPCR. The cell is exposed to a GPCR agonist and then enzymatic activity of the reporter enzyme is detected. Increased reporter enzyme activity indicates that the cell contains a GPCR responsive to the agonist.

The invention is achieved by using ICAST™ protein/protein interaction screening to map signaling pathways. This technology is applicable to a variety of known and unknown GPCRs with diverse functions. They include, but are not limited to, the following sub-families of GPCRs:

5 (a) receptors that bind to amine-like ligands-Acetylcholine muscarinic receptor (M1 to M5), alpha and beta Adrenoceptors, Dopamine receptors (D1, D2, D3 and D4), Histamine receptors (H1 and H2), Octopamine receptor and Serotonin receptors (5HT1, 5HT2, 5HT4, 5HT5, 5HT6, 5HT7);

10 (b) receptors that bind to a peptide ligand-Angiotensin receptor, Bombesin receptor, Bradykinin receptor, C-C chemokine receptors (CCR1 to CCR8, and CCR10), C-X-C type Chemokine receptors (CXC-R5), Cholecystokinin type A receptor, CCK type receptors, Endothelin receptor, Neurotesin receptor, FMLP-related receptors, Somatostatin receptors (type 1 to type 5) and Opioid receptors (type D, K, M, X);

15 (c) receptors that bind to hormone proteins- Follic stimulating hormone receptor, Thyrotrophin receptor and Lutropin-choriogonadotropic hormone receptor;

(d) receptors that bind to neurotransmitters-substance P receptor, Substance K receptor and neuropeptide Y receptor;

(e) Olfactory receptors-Olfactory type 1 to type 11, Gustatory and odorant receptors;

20 (f) Prostanoid receptors-Prostaglandin E2 (EP1 to EP4 subtypes), Prostacyclin and Thromboxane;

(g) receptors that bind to metabotropic substances-Metabotropic glutamate group I to group III receptors;

(h) receptors that respond to physical stimuli, such as light, or to chemical stimuli, such as taste and smell; and

(i) orphan GPCRs-the natural ligand to the receptor is undefined.

ICAST™ provides many benefits to the screening process, including the ability to monitor protein interactions in any sub-cellular compartment-membrane, cytosol and nucleus; the ability to achieve a more physiologically relevant model without requiring protein overexpression; and the ability to achieve a functional assay for receptor binding allowing high information content.

### BRIEF DESCRIPTION OF THE DRAWINGS

FIGURE 1. Cellular expression levels of  $\beta 2$  adrenergic receptor ( $\beta 2AR$ ) and  $\beta$ -arrestin-2 ( $\beta Arr2$ ) in C2 clones. Quantification of  $\beta$ -gal fusion protein was performed using antibodies against  $\beta$ -gal and purified  $\beta$ -gal protein in a titration curve by a standardized ELISA assay. Figure 1A shows expression levels of  $\beta 2AR$ - $\beta gal\Delta\alpha$  clones (in expression vector pICAST ALC). Figure 1B shows expression levels of  $\beta Arr2$ - $\beta gal\Delta\omega$  in expression vector pICAST OMC4 for clones 9-3, -7, -9, -10, -19 and -24, or in expression vector pICAST OMN4 for clones 12-4, -9, -16, -18, -22 and -24.

FIGURE 2. Receptor  $\beta 2AR$  activation was measured by agonist-stimulated cAMP production. C2 cells expressing pICAST ALC  $\beta 2AR$  (clone 5) or parental cells were treated with increasing concentrations of (-)-isoproterenol and 0.1mM IBMX. The quantification of cAMP level was expressed as pmol/well.

FIGURE 3. Interaction of activated receptor  $\beta$ 2AR and arrestin can be measured by  $\beta$ -galactosidase complementation. Figure 3A shows a time course of  $\beta$ -galactosidase activity in response to agonist (-)isoproterenol stimulation in C2 expressing  $\beta$ 2AR- $\beta$ gal $\Delta\alpha$  ( $\beta$ 2AR alone, in expression vector pICAST ALC), or C2 clones, and a pool of C2 co-expressing  $\beta$ 2AR- $\beta$ gal $\Delta\alpha$  and  $\beta$ Arr2- $\beta$ gal $\Delta\omega$  (in expression vectors pICAST ALC and pICAST OMC). Figure 3B shows a time course of  $\beta$  galactosidase activity in response to agonist (-)isoproterenol stimulation in C2 cells expressing  $\beta$ 2AR alone (in expression vector pICAST ALC) and C2 clones co-expressing  $\beta$ 2AR and  $\beta$ Arr1 (in expression vectors ICAST ALC and pICAST OMC).

FIGURE 4. Agonist dose response for interaction of  $\beta$ 2AR and arrestin can be measured by  $\beta$ -galactosidase complementation. Figure 4A shows a dose response to agonists (-)isoproterenol and procaterol in C2 cells co-expressing pICAST ALC  $\beta$ 2AR and pICAST OMC  $\beta$ Arr2 fusion constructs. Figure 4B shows a dose response to agonists (-)isoproterenol and procaterol in C2 cells co-expressing pICAST ALC  $\beta$ 2AR and pICAST OMC  $\beta$ Arr1 fusion constructs.

FIGURE 5. Antagonist mediated inhibition of receptor activity can be measured by  $\beta$ -galactosidase complementation in cells co-expressing  $\beta$ 2AR- $\beta$ gal $\Delta\alpha$  and  $\beta$ Arr- $\beta$ gal $\Delta\omega$ . Figure 5A shows specific inhibition with adrenergic antagonists ICI-118,551 and propranolol of  $\beta$ -galactosidase activity in C2 clones co-expressing pICAST ALC  $\beta$ 2AR and pICAST OMC  $\beta$ Arr2 fusion constructs after incubation with agonist (-)isoproterenol. Figure 5B shows specific inhibition of  $\beta$ -galactosidase activity with adrenergic antagonists ICI-118,551

and propranolol in C2 clones co-expressing pICAST ALC  $\beta$ 2AR and pICAST OMC  $\beta$ Arr1 fusion constructs in the presence of agonist (-)isoproterenol.

FIGURE 6. C2 cells expressing adenosine receptor A2a show cAMP induction in response to agonist (CGC-21680) treatment. C2 parental cells and C2 cells co-expressing pICAST ALC A2aR and pICAST OMC  $\beta$ Arr1 as a pool or as selected clones were measured for agonist-induced cAMP response (pmol/well).

FIGURE 7. Agonist stimulated cAMP response in C2 cells co-expressing Dopamine receptor D1 (D1- $\beta$ gal $\Delta\alpha$ ) and  $\beta$ -arrestin-2 ( $\beta$ Arr2- $\beta$ gal $\Delta\omega$ ). The clone expressing  $\beta$ Arr2- $\beta$ gal $\Delta\omega$  (Arr2 alone) was used as a negative control in the assay. Cells expressing D1- $\beta$ gal $\Delta\alpha$  in addition to  $\beta$ Arr2- $\beta$ gal $\Delta\omega$  responded agonist treatment (3-hydroxytyramine hydrochloride at 3  $\mu$ M). D1(PIC2) or D1(PIC3) designate D1 in expression vector pICAST ALC2 or pICAST ALC4, respectively.

FIGURE 8. Variety of mammalian cell lines can be used to generate stable cells for monitoring GPCR and arrestin interactions. FIGURE 8A, FIGURE 8B and FIGURE 8C show the examples of HEK293, CHO and CHW cell lines co-expressing adrenergic receptor  $\beta$ 2AR and arrestin fusion proteins of  $\beta$ -galactosidase mutants. The  $\beta$ -galactosidase activity was used to monitor agonist-induced interaction of  $\beta$ 2AR and arrestin proteins.

FIGURE 9. Beta-gal complementation can be used to monitor  $\beta$ 2 adrenergic receptor homo-dimerization. FIGURE 9A shows  $\beta$ -galactosidase activity in HEK293 clones co-expressing pICAST ALC  $\beta$ 2AR and pICAST OMC  $\beta$ 2AR. FIGURE 9B shows a cAMP response to agonist (-)isoproterenol in HEK 293 clones co-expressing pICAST ALC  $\beta$ 2AR

and pICAST OMC  $\beta$ 2AR. HEK293 parental cells were included in the assays as negative controls.

FIGURE 10A. pICAST ALC: Vector for expression of  $\beta$ -gal $\Delta\alpha$  as a C-terminal fusion to the target protein. This construct contains the following features: MCS, multiple cloning site for cloning the target protein in frame with the  $\beta$ -gal $\Delta\alpha$ ; GS Linker, (GGGGS)<sub>n</sub>; NeoR, neomycin resistance gene; IRES, internal ribosome entry site; ColE1ori, origin of replication for growth in E. coli; 5'MoMuLV LTR and 3'MoMuLV LTR, viral promotor and polyadenylation signals from the Moloney Murine leukemia virus.

FIGURE 10B. Nucleotide sequence for pICAST ALC.

FIGURE 11A. pICAST ALN: Vector for expression of  $\beta$ -gal $\Delta\alpha$  as an N-terminal fusion to the target protein. This construct contains the following features: MCS, multiple cloning site for cloning the target protein in frame with the  $\beta$ -gal $\Delta\alpha$ ; GS Linker, (GGGGS)<sub>n</sub>; NeoR, neomycin resistance gene; IRES, internal ribosome entry site; ColE1ori, origin of replication for growth in E. coli; 5'MoMuLV LTR and 3'MoMuLV LTR, viral promotor and polyadenylation signals from the Moloney Murine leukemia virus.

FIGURE 11B. Nucleotide sequence for pICAST ALN.

FIGURE 12A. pICAST OMC: Vector for expression of  $\beta$ -gal $\Delta\omega$  as a C-terminal fusion to the target protein. This construct contains the following features: MCS, multiple cloning site for cloning the target protein in frame with the  $\beta$ -gal $\Delta\omega$  ; GS Linker, (GGGGS)<sub>n</sub>; Hygro, hygromycin resistance gene; IRES, internal ribosome entry site; ColE1ori, origin of replication for growth in E. coli; 5'MoMuLV LTR and 3'MoMuLV LTR, viral promotor and polyadenylation signals from the Moloney Murine leukemia virus.

FIGURE 12B. Nucleotide sequence for pICAST OMC.

FIGURE 13A. pICAST OMN: Vector for expression of  $\beta$ -gal $\Delta\omega$  as an N-terminal fusion to the target protein. This construct contains the following features: MCS, multiple cloning site for cloning the target protein in frame with the  $\beta$ -gal $\Delta\omega$ ; GS Linker, (GGGGS)<sub>n</sub>; Hygro, hygromycin resistance gene; IRES, internal ribosome entry site; ColE1ori, origin of replication for growth in E. coli; 5'MoMuLV LTR and 3'MoMuLV LTR, viral promotor and polyadenylation signals from the Moloney Murine leukemia virus.

FIGURE 13B. Nucleotide sequence for pICAST OMN.

FIGURE 14. pICAST ALC  $\beta$ Arr2: Vector for expression of  $\beta$ -gal $\Delta\alpha$  as a C-terminal fusion to  $\beta$ -arrestin-2. The coding sequence of human  $\beta$ -arrestin-2 (Genebank Accession Number: NM\_004313) was cloned in frame to  $\beta$ -gal $\Delta\alpha$  in a pICAST ALC vector.

FIGURE 15. pICAST OMC  $\beta$ Arr2: Vector for expression of  $\beta$ -gal $\Delta\omega$  as a C-terminal fusion to  $\beta$ -arrestin-2. The coding sequence of human  $\beta$ -arrestin-2 (Genebank Accession Number: NM\_004313) was cloned in frame to  $\beta$ -gal $\Delta\omega$  in a pICAST OMC vector.

FIGURE 16. pICAST ALC  $\beta$ Arr1: Vector for expression of  $\beta$ -gal $\Delta\alpha$  as a C-terminal fusion to  $\beta$ -arrestin-1. The coding sequence of human  $\beta$ -arrestin-1 (Genebank Accession Number: NM\_004041) was cloned in frame to  $\beta$ -gal $\Delta\alpha$  in a pICAST ALC vector.

FIGURE 17. pICAST OMC  $\beta$ Arr1: Vector for expression of  $\beta$ -gal $\Delta\omega$  as a C-terminal fusion to  $\beta$ -arrestin-1. The coding sequence of human  $\beta$ -arrestin-1 (Genebank Accession Number: NM\_004041) was cloned in frame to  $\beta$ -gal $\Delta\omega$  in a pICAST OMC vector.

FIGURE 18. pICAST ALC  $\beta$ 2AR: Vector for expression of  $\beta$ -gal $\Delta\alpha$  as a C-terminal fusion to  $\beta$ 2 Adrenergic Receptor. The coding sequence of human  $\beta$ 2 Adrenergic Receptor

(Genebank Accession Number: NM\_000024) was cloned in frame to  $\beta$ -gal $\Delta\alpha$  in a pICAST ALC vector.

FIGURE 19. pICAST OMC  $\beta$ 2AR: Vector for expression of  $\beta$ -gal $\Delta\omega$  as a C-terminal fusion  $\beta$ 2 Adrenergic Receptor. The coding sequence of human  $\beta$ 2 Adrenergic Receptor (Genebank Accession Number: NM\_000024) was cloned in frame to  $\beta$ -gal $\Delta\omega$  in a pICAST OMC vector.

FIGURE 20. pICAST ALC A2aR: Vector for expression of  $\beta$ -gal $\Delta\alpha$  as a C-terminal fusion to Adenosine 2a Receptor. The coding sequence of human Adenosine 2a Receptor (Genebank Accession Number: NM\_000675) was cloned in frame to  $\beta$ -gal $\Delta\alpha$  in a pICAST ALC vector.

FIGURE 21. pICAST OMC A2aR: Vector for expression of  $\beta$ -gal $\Delta\omega$  as a C-terminal fusion to Adenosine 2a Receptor. The coding sequence of human Adenosine 2a Receptor (Genebank Accession Number: NM\_000675) was cloned in frame to  $\beta$ -gal $\Delta\omega$  in a pICAST OMC vector.

FIGURE 22. pICAST ALC D1: Vector for expression of  $\beta$ -gal $\Delta\alpha$  as a C-terminal fusion to Dopamine D1 Receptor. The coding sequence of human Dopamine D1 Receptor (Genebank Accession Number: X58987) was cloned in frame to  $\beta$ -gal $\Delta\alpha$  in a pICAST ALC vector.

FIGURE 23. A schematic depicting the method of the invention, which shows that two inactive mutants that become active when they interact.



## DETAILED DESCRIPTION OF PREFERRED EMBODIMENTS

All literature and patents cited in this disclosure are incorporated herein by reference.

The present invention provides a method to interrogate GPCR function and pathways.

The G-protein-coupled superfamily continues to expand rapidly as new receptors are  
5 discovered through automated sequencing of cDNA libraries or genomic DNA. It is  
estimated that several thousand GPCRs may exist in the human genome, as many as 250  
GPCRs have been cloned and only as few as 150 have been associated with ligands. The  
means by which these, or newly discovered orphan receptors, will be associated with their  
10 cognate ligands and physiological functions represents a major challenge to biological and  
biomedical research. The identification of an orphan receptor generally requires an  
individualized assay and a guess as to its function. The interrogation of a GPCR's signaling  
behavior by introducing a replacement receptor eliminates these prerequisites because it can  
be performed with and without prior knowledge of other signaling events. It is sensitive,  
rapid and easily performed and should be applicable to nearly all GPCRs because the  
15 majority of these receptors should desensitize by a common mechanism.

Various approaches have been used to monitor intracellular activity in response to a  
stimulant, e.g., enzyme-linked immunosorbent assay (ELISA); Fluorescence Imaging Plate  
Reader assay (FLIPR™, Molecular Devices Corp., Sunnyvale, CA); EVOscreen™,  
EVOTEC™, Evotec Biosystems GmbH, Hamburg, Germany; and techniques developed by  
20 CELLOMICS™, Cellomics, Inc., Pittsburgh, PA.

Germino, F.J., et al., "Screening for in vivo protein-protein interactions." Proc. Natl.  
Acad. Sci., 90(3): 933-7 (1993), discloses an *in vivo* approach for the isolation of proteins  
interacting with a protein of interest.

Phizicky, E.M., et al., "Protein-protein interactions: methods for detection and analysis." Microbiol. Rev., 59(1): 94-123 (1995), discloses a review of biochemical, molecular biological and genetic methods used to study protein-protein interactions.

Offermanns, et al., " $G\alpha_{15}$  and  $G\alpha_{16}$  Couple a Wide Variety of Receptors to Phospholipase C." J. Biol. Chem., 270(25):15175-80 (1995), discloses that  $G\alpha_{15}$  and  $G\alpha_{16}$  can be activated by a wide variety of G-protein-coupled receptors. The selective coupling of an activated receptor to a distinct pattern of G-proteins is regarded as an important requirement to achieve accurate signal transduction. Id.

Barak et al., "A  $\beta$ -arrestin/Green Fluorescent Protein Biosensor for Detecting G Protein-coupled Receptor Activation." J. Biol. Chem., 272(44):27497-500 (1997) and U.S. Patent No. 5,891,646, disclose the use of a  $\beta$ -arrestin/green fluorescent fusion protein (GFP) to monitor protein translocation upon stimulation of GPCR.

The present invention involves a method for monitoring protein-protein interactions in GPCR pathways as a complete assay using ICAST™ (Intercistronic Complementation Analysis Screening Technology as disclosed in pending U.S. patent application serial no. 053,164, filed April 1, 1998, the entire contents of which are incorporated herein by reference). This invention enables an array of assays, including GPCR binding assays, to be achieved directly within the cellular environment in a rapid, non-radioactive assay format amenable to high-throughput screening. Using existing technology, assays of this type are currently performed in a non-cellular environment and require the use of radioisotopes.

The present invention combined with Tropix ICAST™ and Advanced Discovery Sciences™ technologies, e.g., ultra high-throughput screening, provide highly sensitive cell-based methods for interrogating GPCR pathways which are amendable to high-throughput

screening (HTS). These methods are an advancement over the invention disclosed in U.S. Patent 5,891,646, which relies on microscopic imaging of GPCR components as fusion with Green-fluorescent-protein. Imaging techniques are limited by low-throughput, lack of thorough quantification and low signal to noise ratios. Unlike yeast-based-2-hybrid assays used to monitor protein/protein interactions in high-throughput assays, the present invention is applicable to a variety of cells including mammalian cells, plant cells, protozoa cells such as *E. coli* and cells of invertebrate origin such as yeast, slime mold (*Dictyostelium*) and insects; detects interactions at the site of the receptor target or downstream target proteins rather than in the nucleus; and does not rely on indirect read-outs such as transcriptional activation. The present invention provides assays with greater physiological relevance and fewer false negatives.

Advanced Discovery Sciences™ is in the business of offering custom-developed screening assays optimized for individual assay requirements and validated for automation. These assays are designed by HTS experts to deliver superior assay performance. Advanced Discovery Sciences'™ custom assay development service encompasses the design, development, optimization and transfer of high performance screening assays. Advanced Discovery Sciences™ works to design new assays or convert existing assays to ultra-sensitive luminescent assays ready for the rigors of HTS. Among some of the technologies developed by Advanced Discovery Sciences™ are the cAMP-Screen™ immunoassay system. This system provides ultrasensitive determination of cAMP levels in cell lysates. The cAMP-Screen™ assay utilizes the high-sensitivity chemiluminescent alkaline phosphatase (AP) substrate CSPD® with Sapphire-II™ luminescence enhancer.

### EXAMPLE:

GPCR activation can be measured through monitoring the binding of ligand-activated GPCR by an arrestin. In this assay system, a GPCR, e.g.  $\beta$  adrenergic receptor ( $\beta$  2AR) and a  $\beta$  arrestin are co-expressed in the same cell as fusion proteins with  $\beta$  gal mutants. As illustrated in Figure 1, the  $\beta$ 2AR is expressed as a fusion protein with  $\Delta\alpha$  form of  $\beta$  gal mutant ( $\beta$ 2ADR $\Delta\alpha$ ) and the  $\beta$  arrestin as a fusion protein with the  $\Delta\omega$  mutant of  $\beta$  gal ( $\beta$ -Arr $\Delta\omega$ ). The two fusion proteins exist inside of a resting (or un-stimulated) cell in separate compartments, i.e. membrane for GPCR and cytosol for arrestin, and they can not form an active  $\beta$  galactosidase enzyme. When such a cell is treated with an agonist or a ligand, the ligand-occupied and activated receptor will become a high affinity binding site for Arrestin. The interaction between an activated  $\beta$ 2ADR $\Delta\alpha$  and  $\beta$ -Arr $\Delta\omega$  drives the  $\beta$  gal mutant complementation. The enzyme activity can be measured by using an enzyme substrate, which upon cleavage releases a product measurable by colorimetry, fluorescence, chemiluminescence (e.g. Tropic product GalScreen<sup>TM</sup>).

### Experiment protocol-

1. In the first step, the expression vectors for  $\beta$ 2ADR $\Delta\alpha$  and  $\beta$ Arr2 $\Delta\omega$  were engineered in selectable retroviral vectors pICAST ALC, as described in Figure 18 and pICAST OMC, as in Figure 15.

2. In the second step, the two expression constructs were transduced into either C2C12 myoblast cells, or other mammalian cell lines, such as COS-7, CHO, A431, HEK 293, and CHW. Following selection with antibiotic drugs, stable clones expressing both fusion

proteins at appropriate levels were selected.

3. In the last step, the cells expressing both  $\beta 2\text{ADR}\Delta\alpha$  and  $\beta\text{Arr}2\Delta\omega$  were tested for response by agonist/ligand stimulated  $\beta$  galactosidase activity. Triplicate samples of cells were plated at 10,000 cells in 100 microliter volume into a well of 96-well culture plate. Cells were cultured for 24 hours before assay. For agonist assay (Figure 3 and 4), cells were treated with variable concentrations of agonist, for example, (-) isoproterenol, procaterol, dobutamine, terbutiline or L-L-phenylephrine for 60 min at 37 C. The induced  $\beta$  galactosidase activity was measured by addition of Tropix GalScreen<sup>TM</sup> substrate (Applied Biosystems) and luminescence measured in a Tropix TR717<sup>TM</sup> luminometer (Applied Biosystems). For antagonist assay (Figure 5), cells were pre-incubated for 10 min in fresh medium without serum in the presence of ICI-118,551 or propranolol followed by addition of 10 micro molar (-) isoproterenol.

The assays of this invention, and their application and preparation have been described both generically, and by specific example. The examples are not intended as limiting. Other substituent identities, characteristics and assays will occur to those of ordinary skill in the art, without the exercise of inventive faculty. Such modifications remain within the scope of the invention, unless excluded by the express recitation of the claims advanced below.

**WHAT IS CLAIMED IS:**

1. A method of assessing the effect of a test condition on G-protein-coupled receptor (GPCR) pathway activity, comprising:

a) providing a cell that expresses a GPCR as a fusion protein to one mutant form of reporter enzyme and an interacting protein partner as a fusion to another mutant form of enzyme;

b) exposing the cell to a ligand for said GPCR under said test condition; and

c) monitoring activation of said GPCR by complementation of said reporter enzyme;

wherein increased reporter enzyme activity in the cell compared to that which occurs in the absence of said test condition indicates increased GPCR interaction with its interacting protein partner compared to that which occurs in the absence of said test condition, and decreased reporter enzyme activity in the cell compared to that which occurs in the absence of said test condition indicates decreased GPCR interaction with its interacting protein partner compared to that which occurs in the absence of said test condition.

2. A method according to Claim 1, wherein the test condition is the presence in the cell of a kinase.

3. A method according to Claim 1, wherein the test condition is the presence in the cell of a G-protein.

4. A method according to Claim 1, wherein the test condition is the exposure of the cell to a compound selected from GPCR agonists and GPCR antagonists.

5. A method according to Claim 1, wherein the test condition is co-expression in the cell of a second receptor.

6. A method according to Claim 5, wherein the second receptor is a GPCR receptor.

7. A method according to Claim 5, wherein homo-dimerization of GPCR is determined.

8. A method according to Claim 5, wherein hetero-dimerization of GPCR is determined.

5 9. A method for screening a  $\beta$ -arrestin protein or an unidentified arrestin or arrestin-like protein or fragment and mutant form thereof for the ability to bind to activated GPCRs, comprising:

a) providing a cell that:

i) expresses at least one GPCR as a fusion protein to a reporter enzyme; and

10 ii) contains a conjugate comprising a test  $\beta$ -arrestin protein as a fusion protein with another reporter enzyme;

b) exposing the cell to a ligand for said at least one GPCR; and

c) detecting enzymatic activity of the complemented reporter enzyme;

wherein an increase in enzymatic activity in the cell indicates  $\beta$ -arrestin protein

15 binding to the activated GPCR.

10. A method for screening a test compound for G-protein-coupled receptor (GPCR) agonist activity, comprising:

a) providing a cell that expresses a GPCR as a fusion protein to one mutant form of reporter enzyme and an arrestin protein as a fusion to another mutant form of enzyme;

20 b) exposing the cell to a test compound; and

c) detecting complementation of said reporter enzyme;

wherein increased reporter enzyme activity after exposure of the cell to the test compound indicates GPCR agonist activity of the test compound.

11. A method according to Claim 10, wherein the cell expresses a GPCR whose function is known.

12. A method according to Claim 10, wherein the cell expresses a GPCR whose function is unknown.

5 13. A method according to Claim 10, wherein the cell expresses an odorant or taste GPCR.

14. A method according to Claim 10, wherein the cell expresses a GPCR a  $\beta$ -adrenergic GPCR.

10 15. A method according to Claim 10, wherein the cell is selected from the group consisting of mammalian cells, cells of invertebrate origin, plant cells and protozoa cells.

16. A method according to Claim 10, wherein the cell endogenously expresses a GPCR.

17. A method according to Claim 10, wherein the cell has been transformed to express a GPCR not endogenously expressed by such a cell.

15 18. A method of screening a test compound for G-protein-coupled receptor (GPCR) antagonist activity, comprising:

a) providing a cell that expresses a GPCR as a fusion protein to one mutant form of reporter enzyme and an arrestin protein as a fusion to another mutant form of enzyme;

b) exposing the cell to said test compound;

20 c) exposing the cell to an agonist for said GPCR; and

d) detecting complementation of said reporter enzyme;

where exposure to the agonist occurs at the same time as, or subsequent to, exposure to the test compound, and wherein decreased reporter enzyme activity after exposure of the



cell to the test compound indicates that the test compound is an antagonist for said GPCR.

19. A method of screening a cell for the presence of a G-protein-coupled receptor (GPCR) responsive to a GPCR agonist, comprising:

- a) providing a cell, said cell containing a conjugate comprising a  $\beta$ -arrestin protein as a fusion protein with a reporter enzyme;
- b) exposing the cell to a GPCR agonist; and
- c) detecting enzymatic activity of the reporter enzyme;

wherein an increase in enzymatic activity after exposure of the cell to the GPCR agonist indicates that the cell contains a GPCR responsive to said agonist.

20. A method of screening a plurality of cells for those cells which contain a G-protein-coupled receptor (GPCR) responsive to a GPCR agonist, comprising:

- a) providing a plurality of cells, said cells containing a conjugate comprising a  $\beta$ -arrestin protein as a fusion protein with a reporter enzyme;
- b) exposing the cells to a GPCR agonist; and
- c) detecting enzymatic activity of the reporter enzyme;

wherein an increase in enzymatic activity after exposure to the GPCR agonist indicates  $\beta$ -arrestin protein binding to a GPCR, thereby indicating that the cell contains a GPCR responsive to said GPCR agonist.

21. A method according to Claim 20, wherein the plurality of cells are contained in a tissue.

22. A method according to Claim 20, wherein the plurality of cells are contained in an organ.

23. A method according to Claim 20, wherein step (b) comprises exposing the cells to a plurality of GPCR agonists or ligand libraries.

24. A substrate having deposited thereon a plurality of cells, said cells expressing at least one GPCR as a fusion protein to one mutant form of reporter enzyme and an arrestin protein as a fusion to another mutant form of enzyme.

25. A substrate according to Claim 24, wherein the substrate contains an enzyme-labile chemical group which, upon cleavage by the reporter enzyme, releases a product measurable by colorimetry, fluorescence or chemiluminescence.

26. A substrate according to Claim 24, wherein the substrate is made of a material selected from glass, plastic, ceramic, semiconductor, silica, fiber optic, diamond, biocompatible monomer and biocompatible polymer materials.

27. A method of detecting G-protein-coupled receptor (GPCR) pathway activity in a cell expressing at least one GPCR and containing  $\beta$ -arrestin protein as a fusion protein with a reporter enzyme; wherein said enzymatic activity indicates activation of the GPCR pathway.

28. A method according to Claim 27, where the cells are deposited on a substrate prior to detecting said enzymatic activity.

29. A method according to Claim 27, wherein said cell is contained in a tissue.

30. A method according to Claim 27, wherein said cell is contained in an organ.

## ABSTRACT

Methods for detecting G-protein coupled receptor (GPCR) activity; methods of assaying GPCR activity; and methods of screening for GPCR ligands, G-protein-coupled receptor kinase (GRK) activity, and compounds that interact with components of the GPCR regulatory process are described.

Cellular Expression of  $\beta_2$ AR- $\beta$ gal $\Delta\alpha$  Fusion Protein in C2 Clones  
(measured by anti- $\beta$ -gal ELISA)

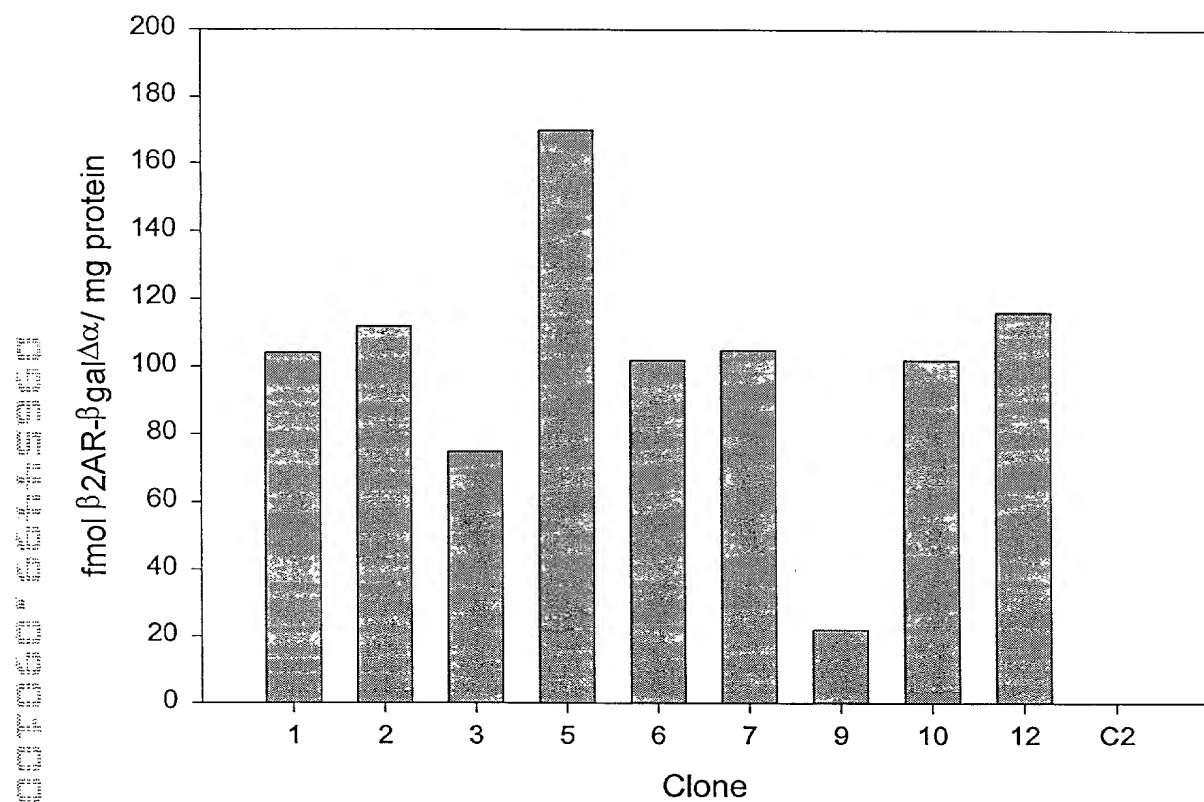


FIGURE 1A

Cellular expression of  $\beta$ Arr2- $\beta$ gal $\Delta\omega$  fusion protein in C2 clones  
(measured by anti- $\beta$  gal ELISA)

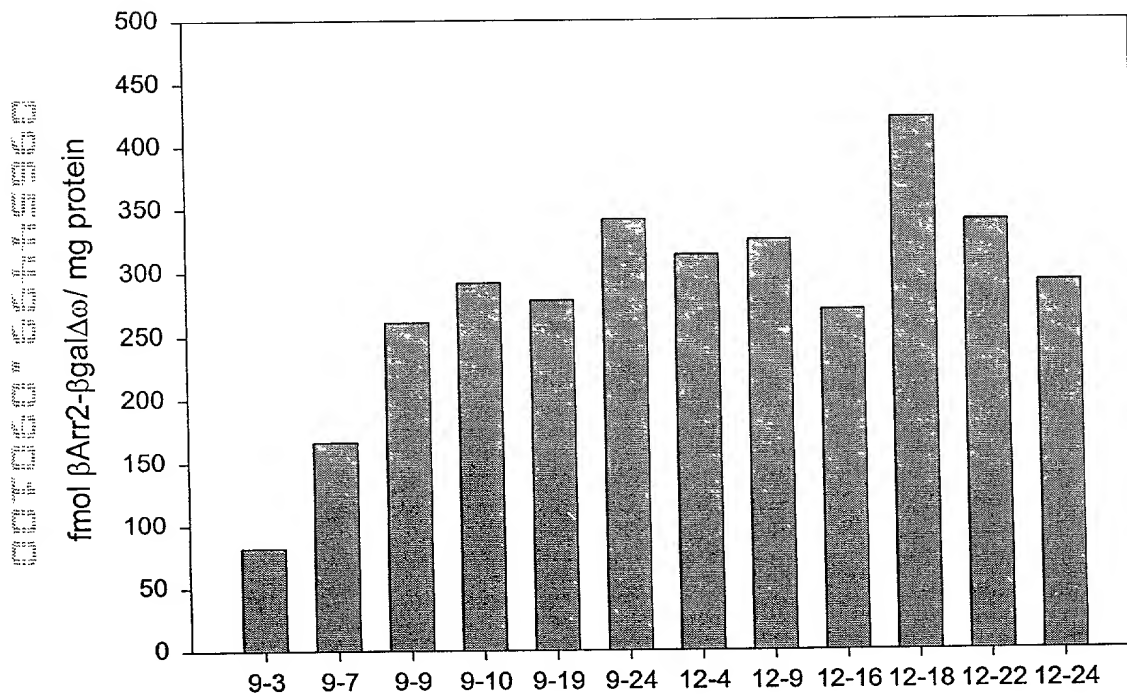


FIGURE 1B

# Agonist Stimulated cAMP Response in C2 Cells Expressing $\beta 2AR$ - $\beta gal\Delta\alpha$

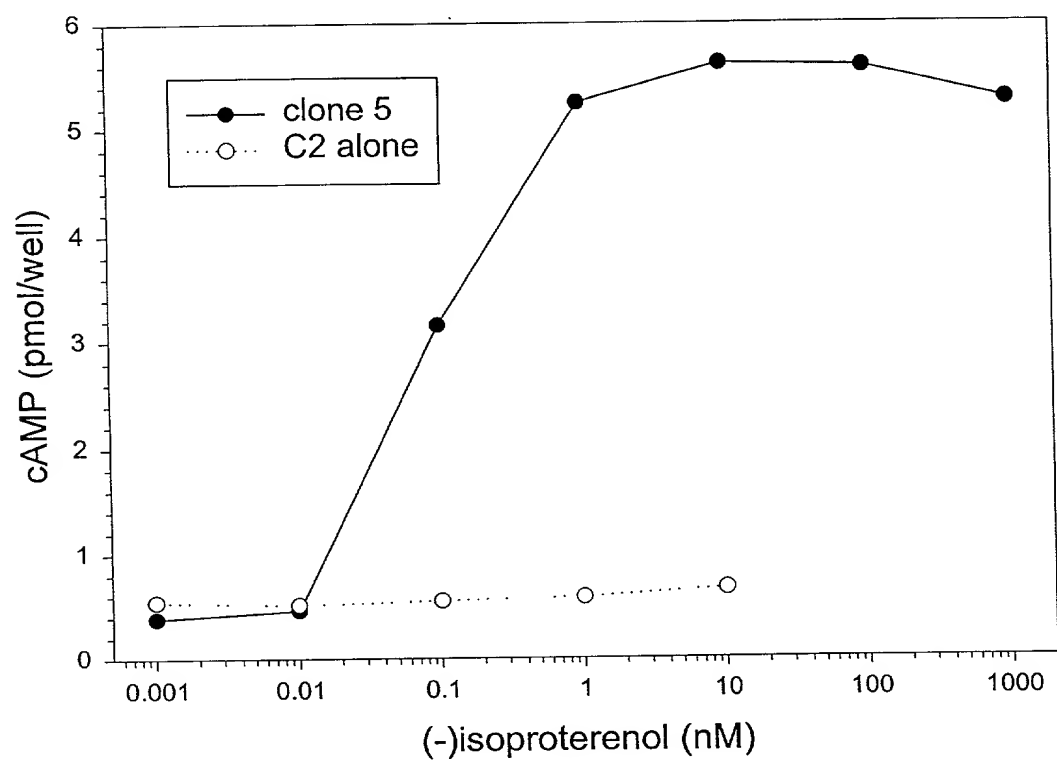


FIGURE 2

$\beta$ -galactosidase Complementation as a Measurement for  $\beta$ 2AR- $\beta$ gal $\Delta\alpha$  interacting with  $\beta$ Arrestin2- $\beta$ gal $\Delta\omega$  upon agonist Stimulation

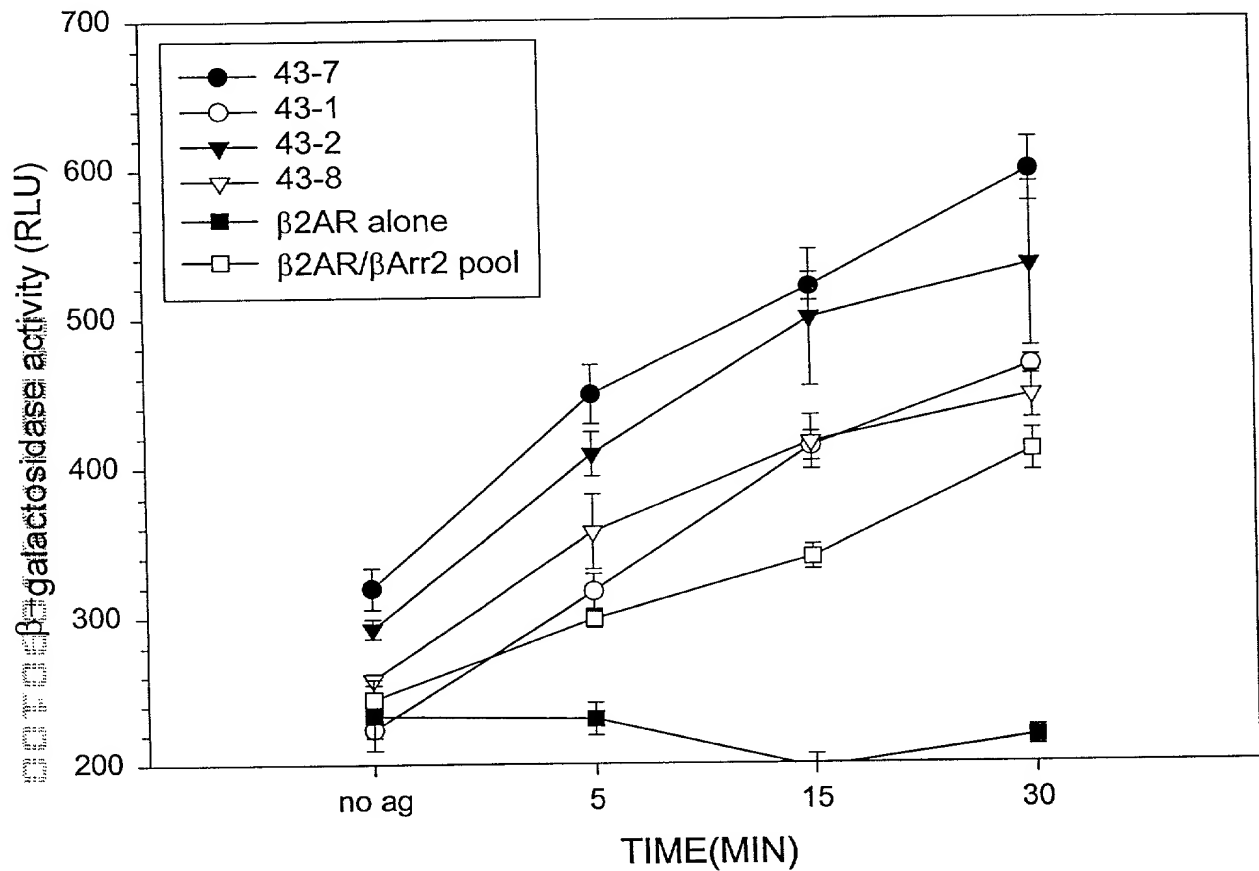


FIGURE 3A

$\beta$ -galactosidase Complementation as a Measurement for  $\beta$ 2AR- $\beta$ gal $\Delta\alpha$  Interaction with  $\beta$ Arrestin1- $\beta$ gal $\Delta\omega$  upon Agonist Stimulation

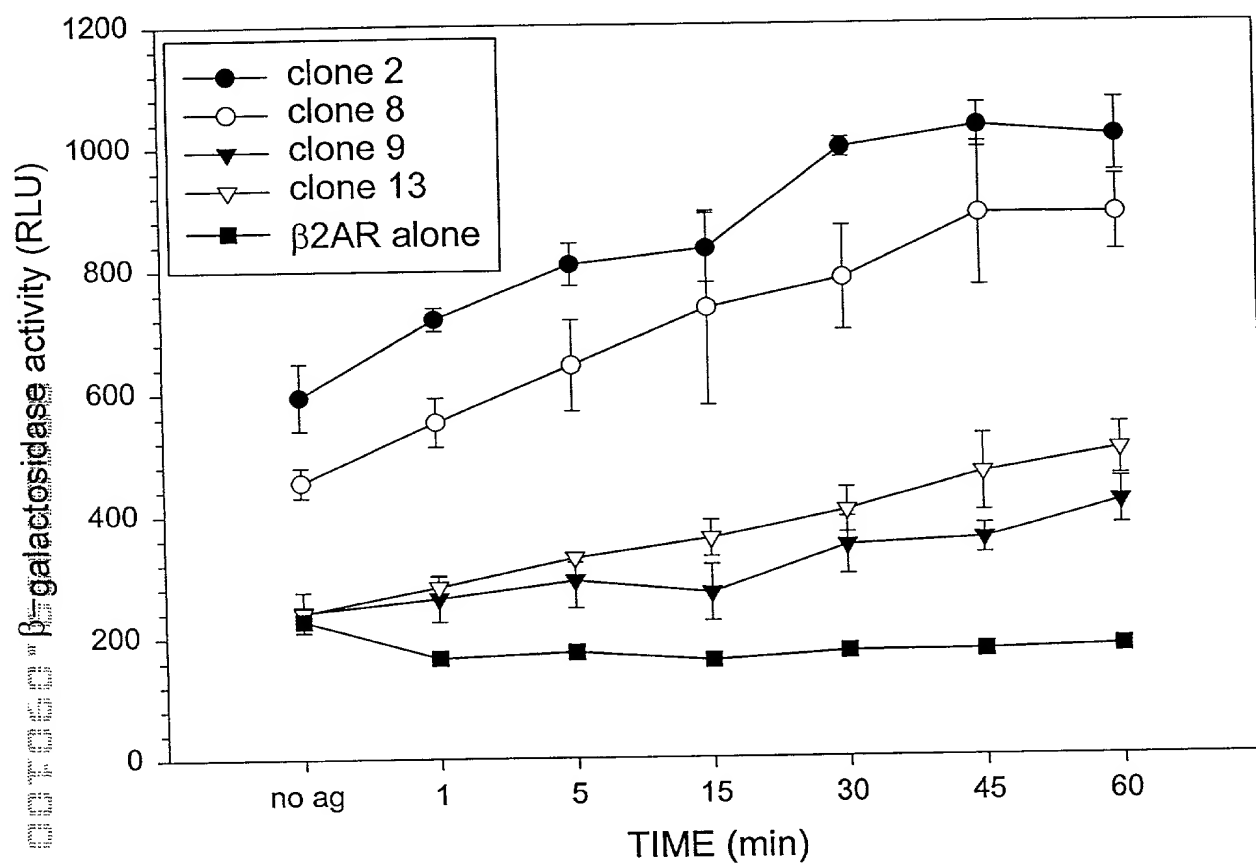


FIGURE 3B



$\beta$ -galactosidase Activity in Response to Agonist in C2 Cells  
Coexpressing  $\beta$ 2AR- $\beta$ gal $\Delta\alpha$  and  $\beta$ Arrestin2- $\beta$ gal $\Delta\omega$  Fusion Proteins

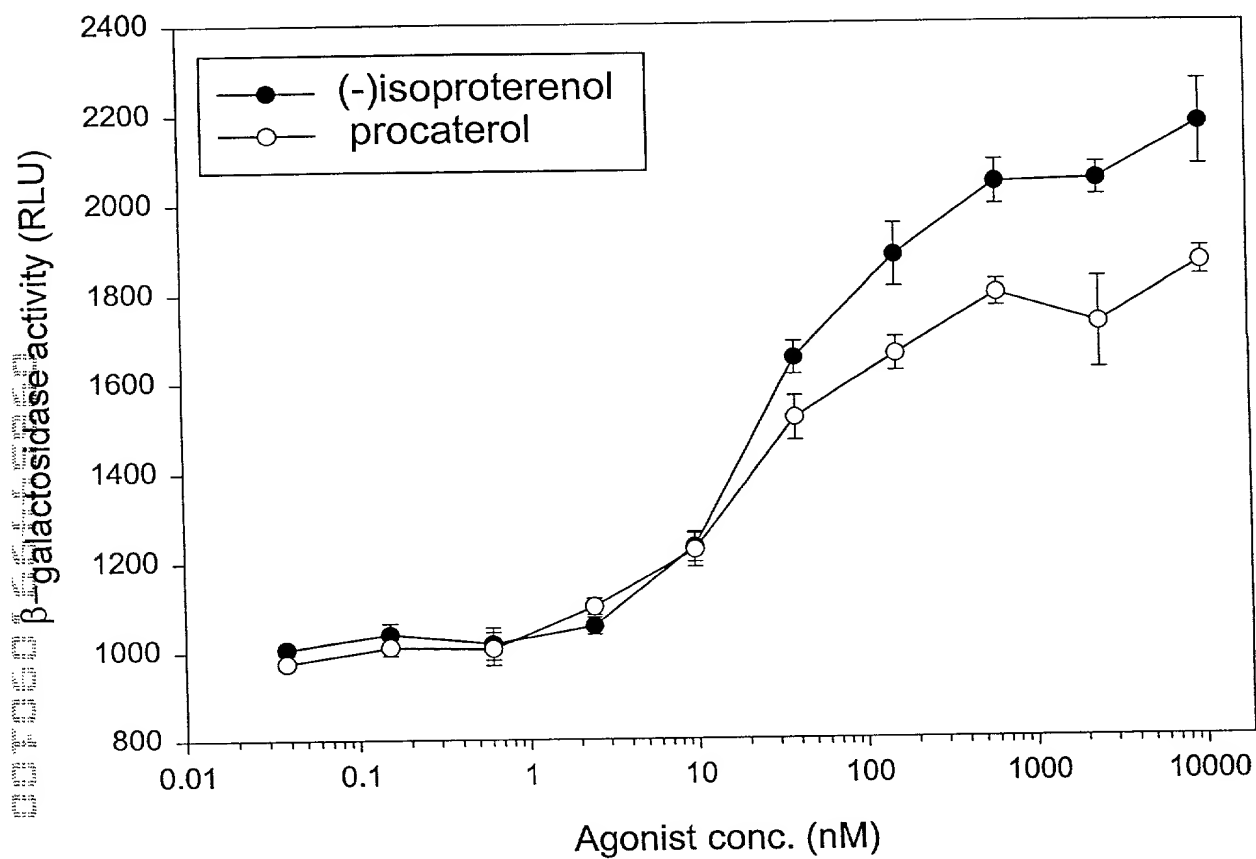


FIGURE 4A

$\beta$ -galactosidase Activity in Response to Agonist in C2 Cells  
Coexpressing  $\beta$ 2AR- $\beta$ gal $\Delta\alpha$  and  $\beta$ Arrestin1- $\beta$ gal $\Delta\omega$  Fusion Proteins

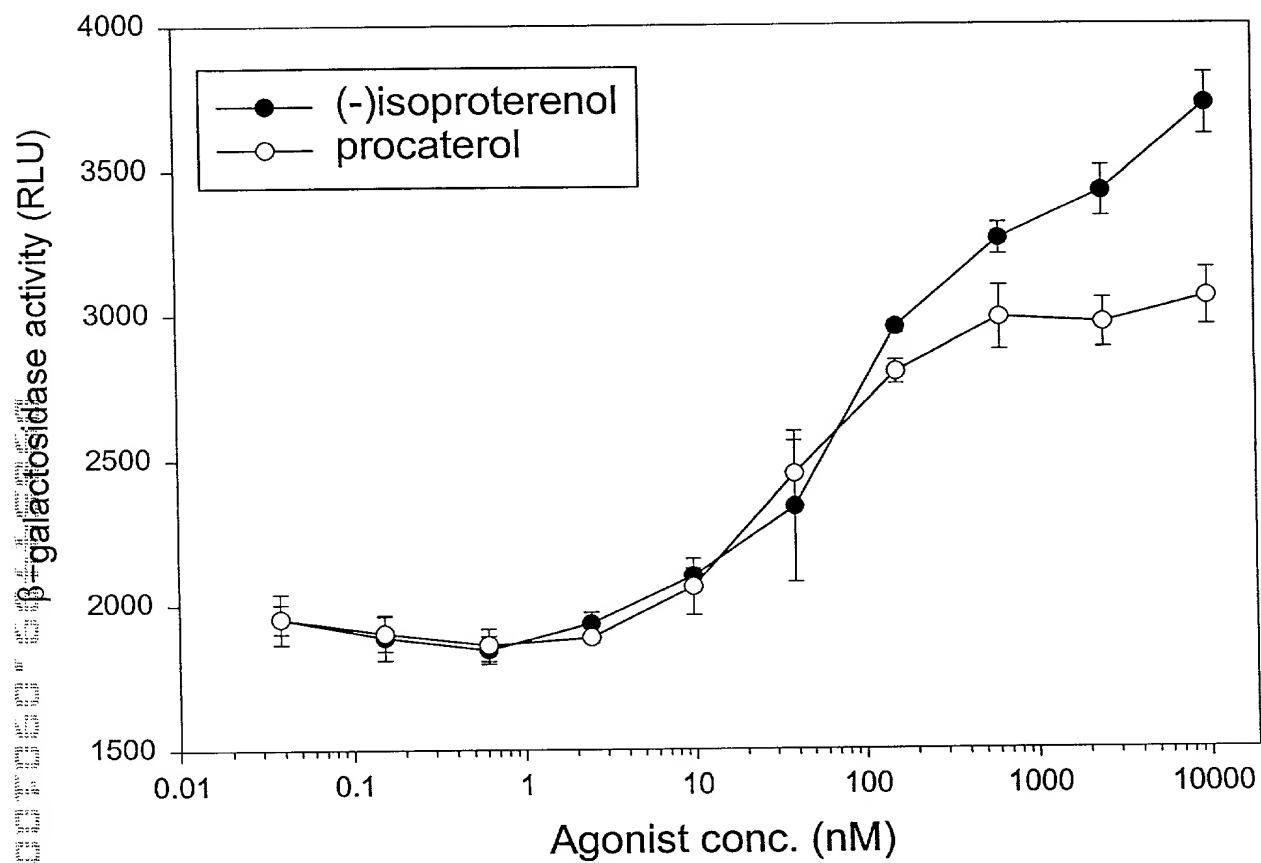


FIGURE 4B

Inhibition of  $\beta$ -galactosidase activity in C2 Cells Coexpressing  $\beta$ 2AR- $\beta$ gal $\Delta\alpha$  and  $\beta$ Arrestin2- $\beta$ gal $\Delta\omega$  Fusion Proteins

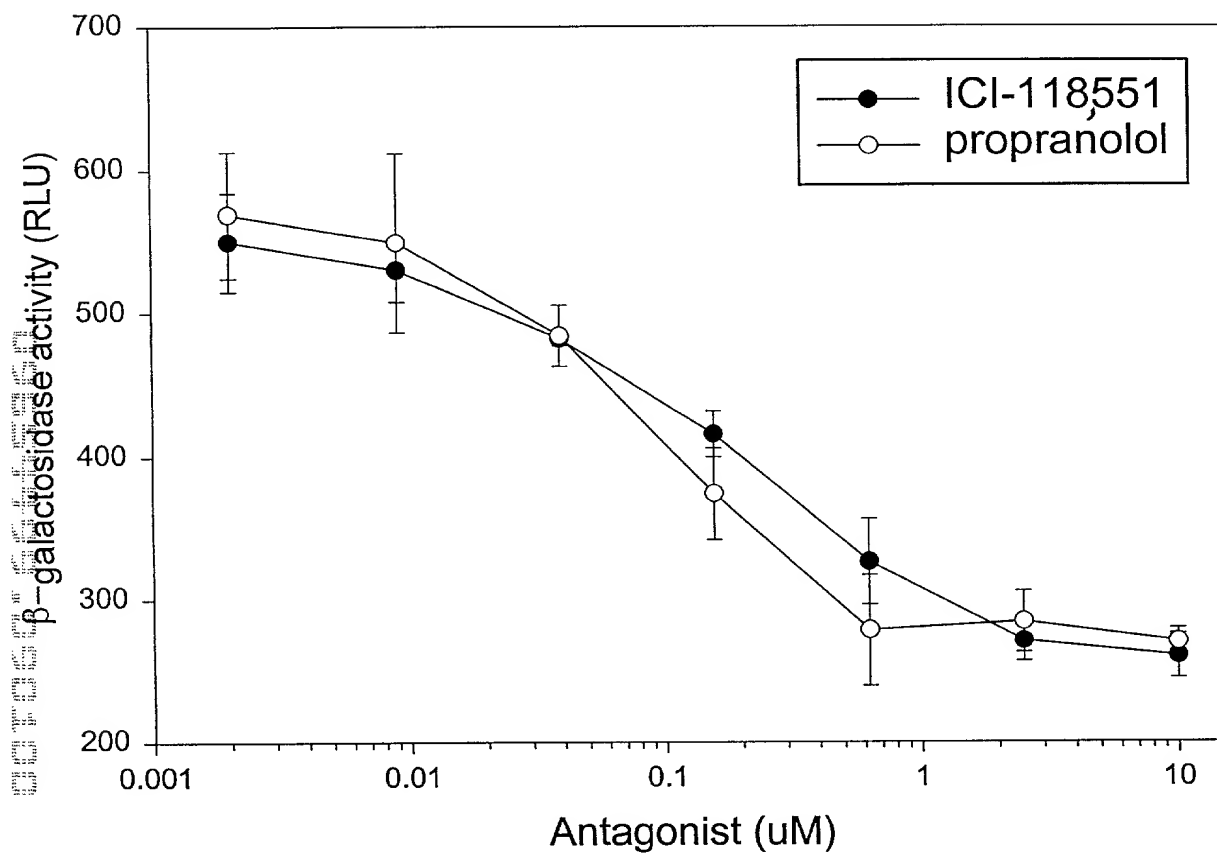


FIGURE 5A

Antagonist Inhibition of  $\beta$ -galactosidase Activity in C2 Cells  
Coexpressing  $\beta$ 2AR- $\beta$ gal $\Delta\alpha$  and  $\beta$ Arrestin1- $\beta$ gal $\Delta\omega$  Fusion Proteins

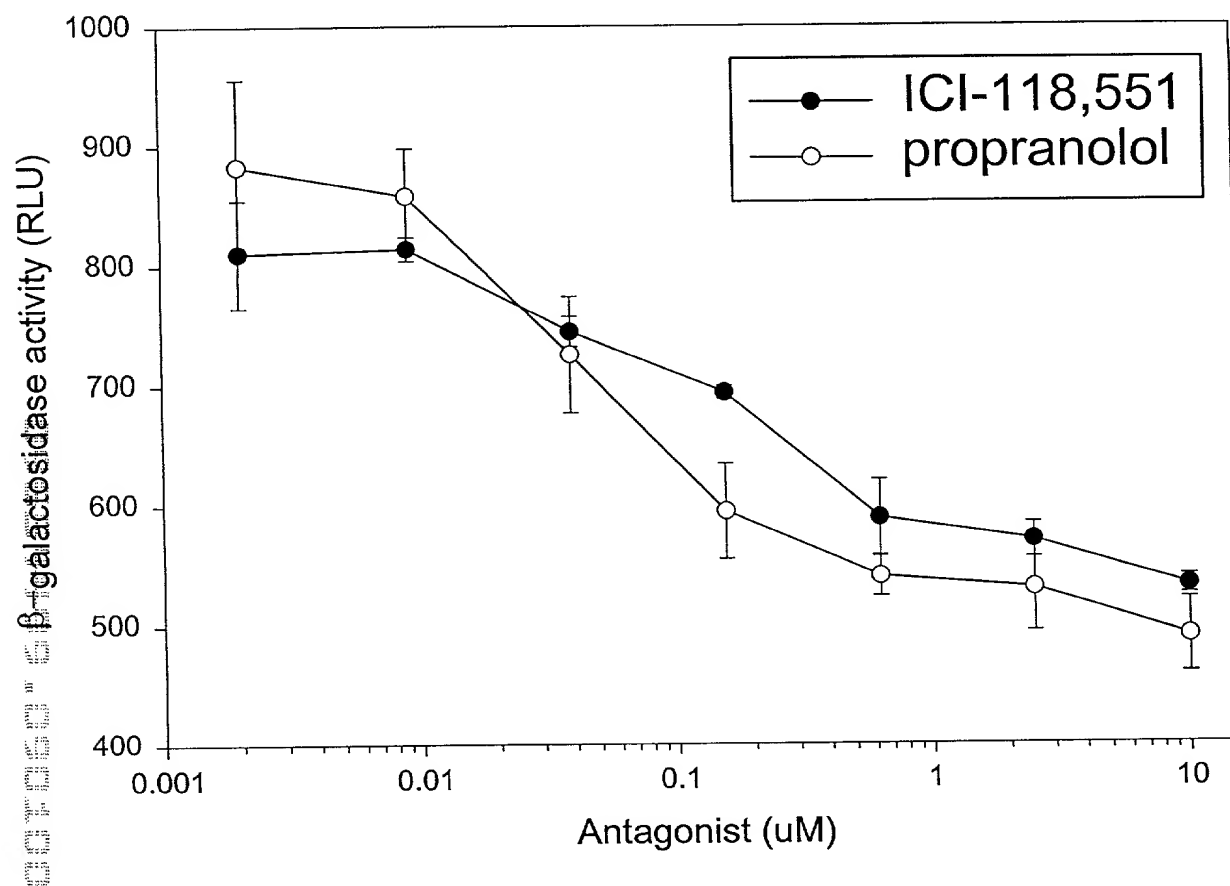


Figure 5B

Agonist Stimulated cAMP Response in Clones or Pools of C2 Cells Coexpressing A2aR-βgalΔα and βArrestin1-βgalΔω Fusion Proteins

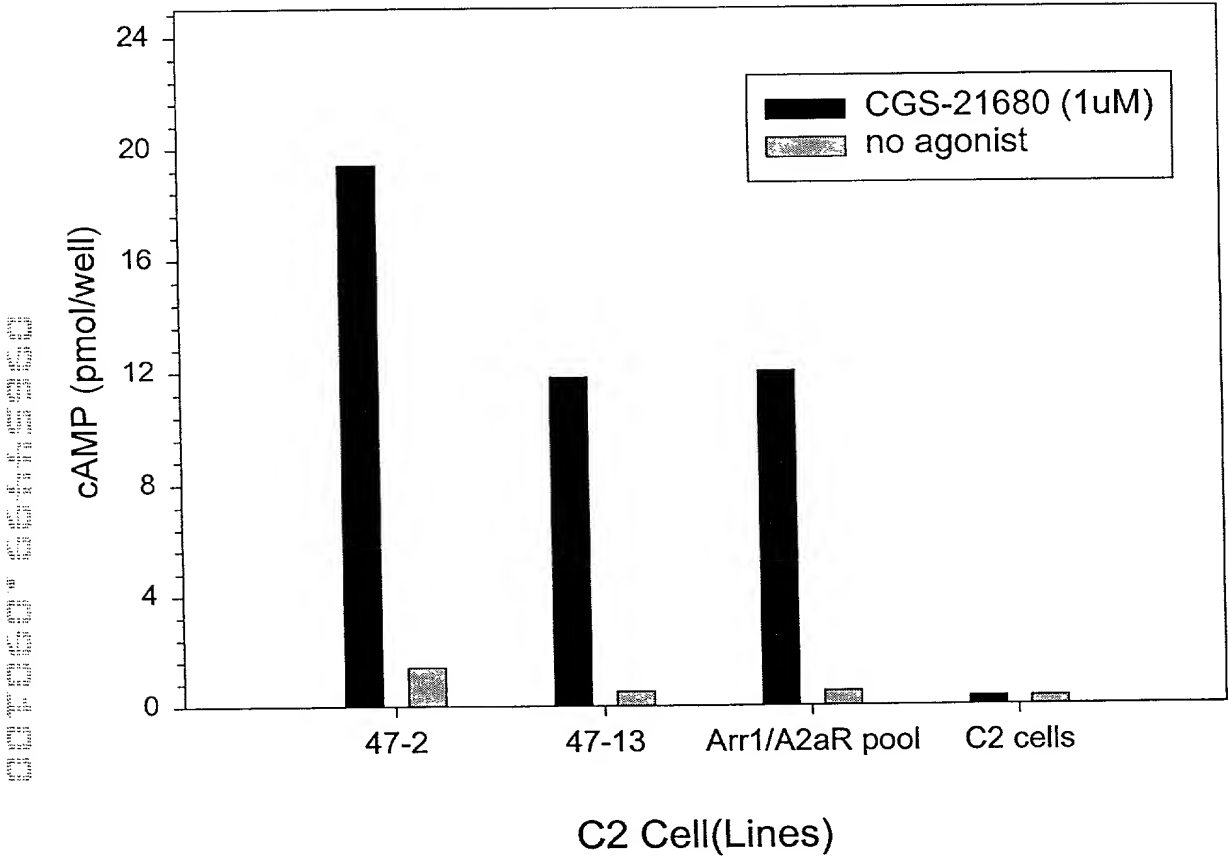


FIGURE 6

# Agonist Stimulated cAMP Response in Clones or Pools of C2 Cells Expressing D1- $\beta$ gal $\Delta\alpha$ and $\beta$ Arrestin2- $\beta$ gal $\Delta\omega$ Fusion Proteins

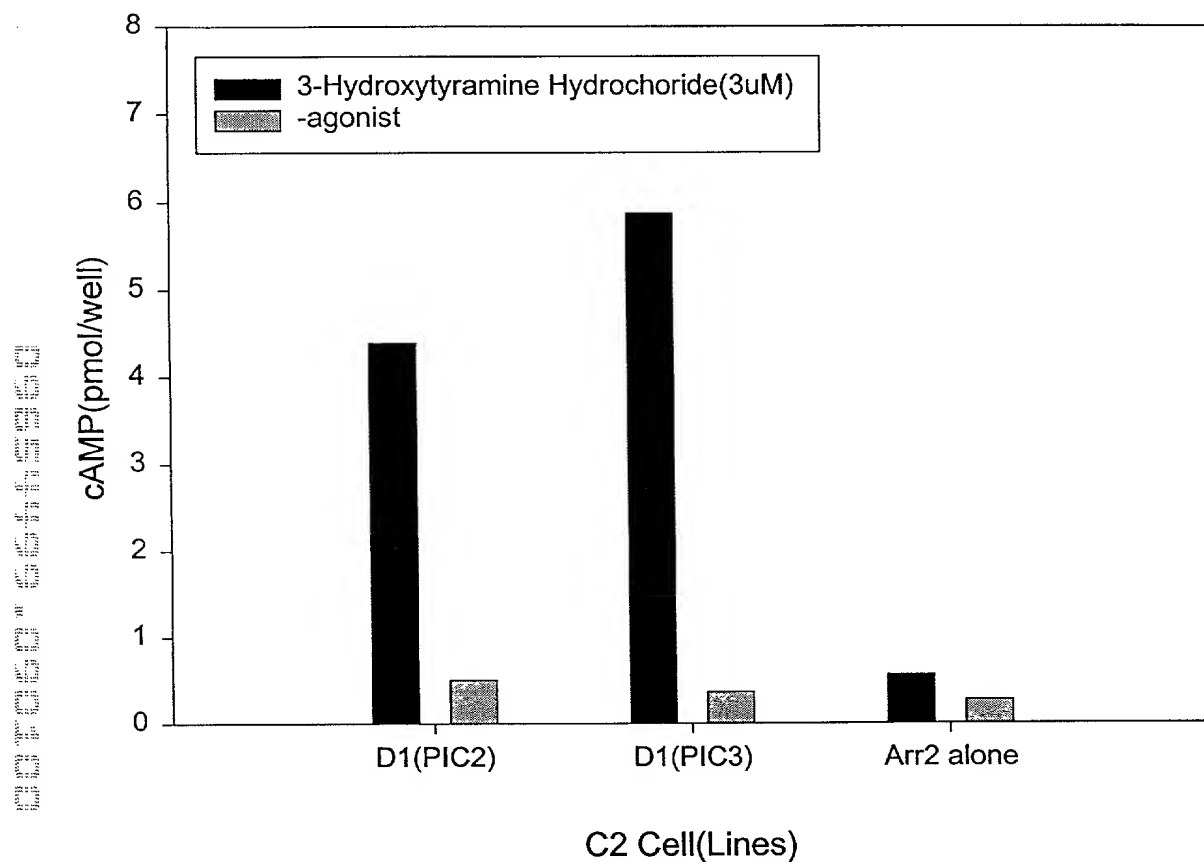


FIGURE 7

**$\beta_2$ AR- $\beta$ gal $\Delta\omega$  and  $\beta$ arr2- $\beta$ gal $\Delta\alpha$  Interaction in HEK293  
Clones in Response to Isoproterenol Treatment (1  $\mu$ M)**

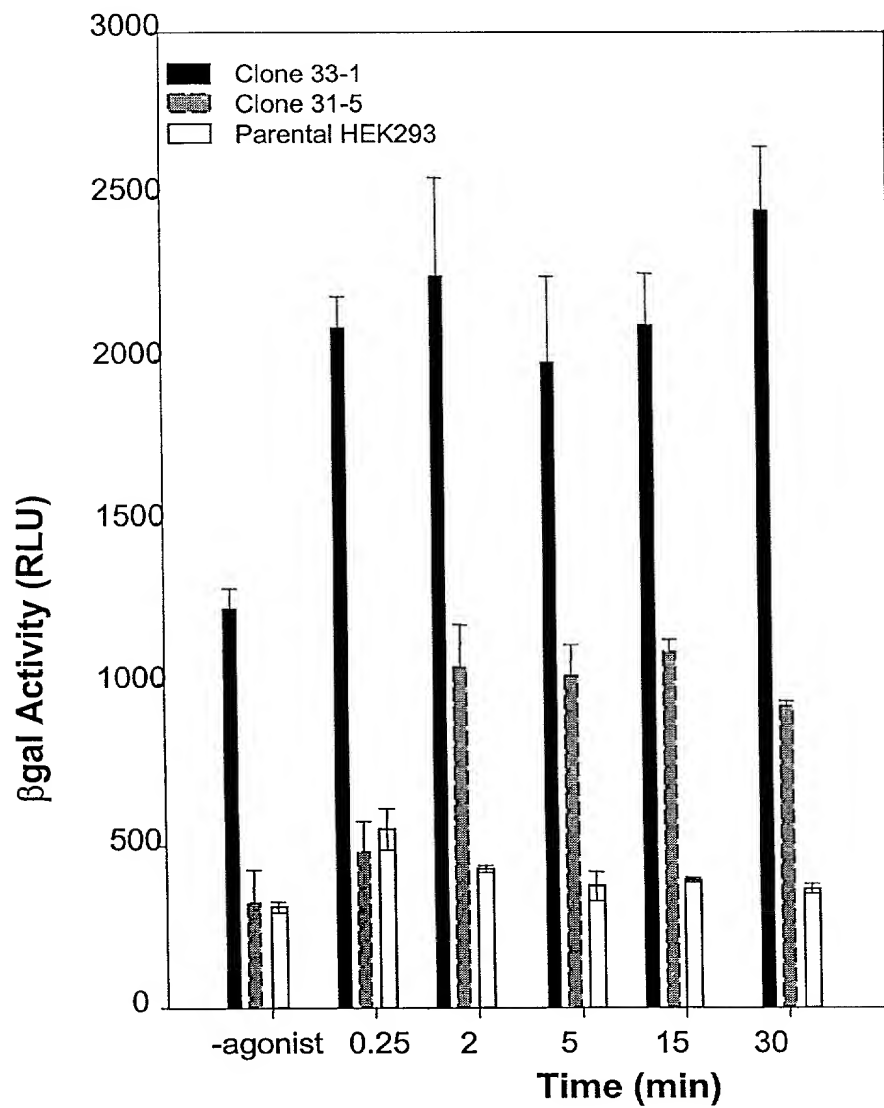


FIGURE 8A

$\beta$ 2AR- $\beta$ gal $\Delta\alpha$  and  $\beta$ Arr1- $\beta$ gal $\Delta$  Interaction in a CHO Pool  
in Response to Isoproterenol Treatment(10uM)

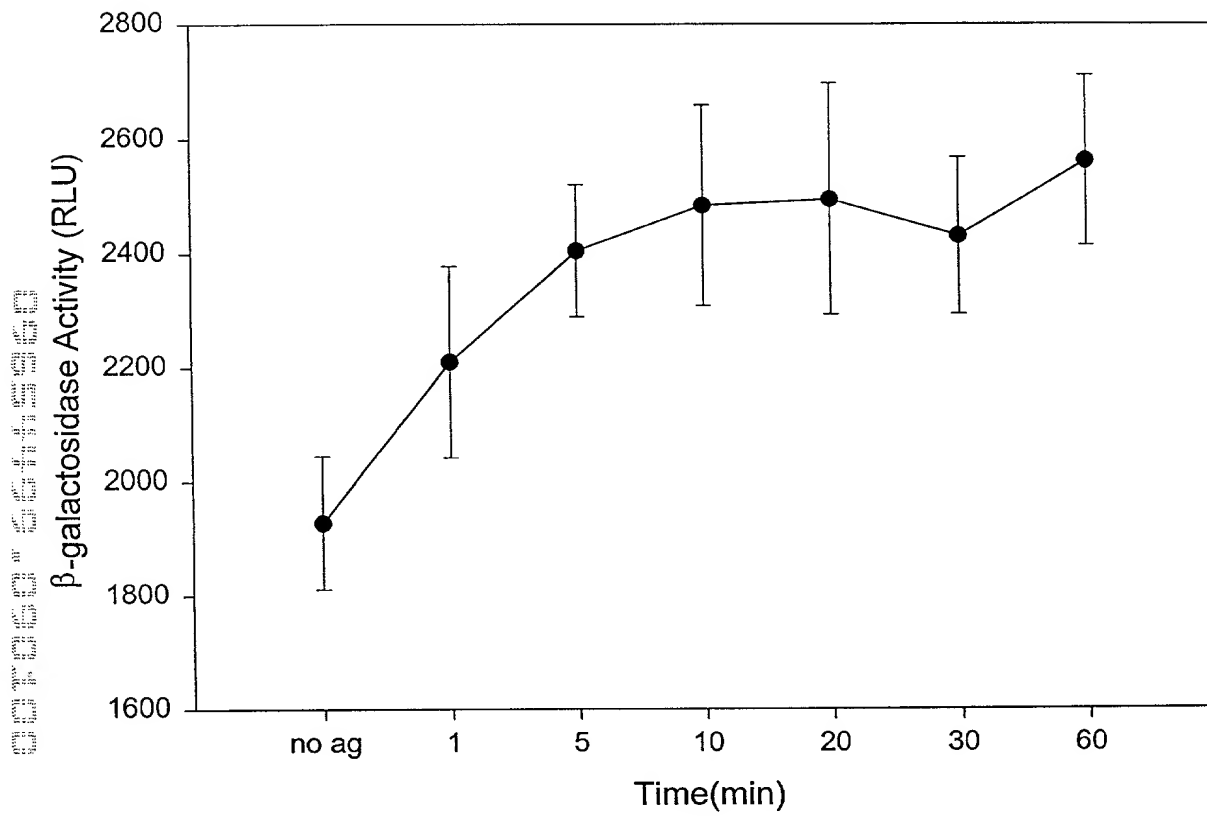


FIGURE 8B



$\beta$ 2AR- $\beta$ gal $\Delta\alpha$  and  $\beta$ Arr2- $\beta$ gal $\Delta\omega$  Interaction in CHW Clone  
in Response to Isoproterenol Treatment (10uM)

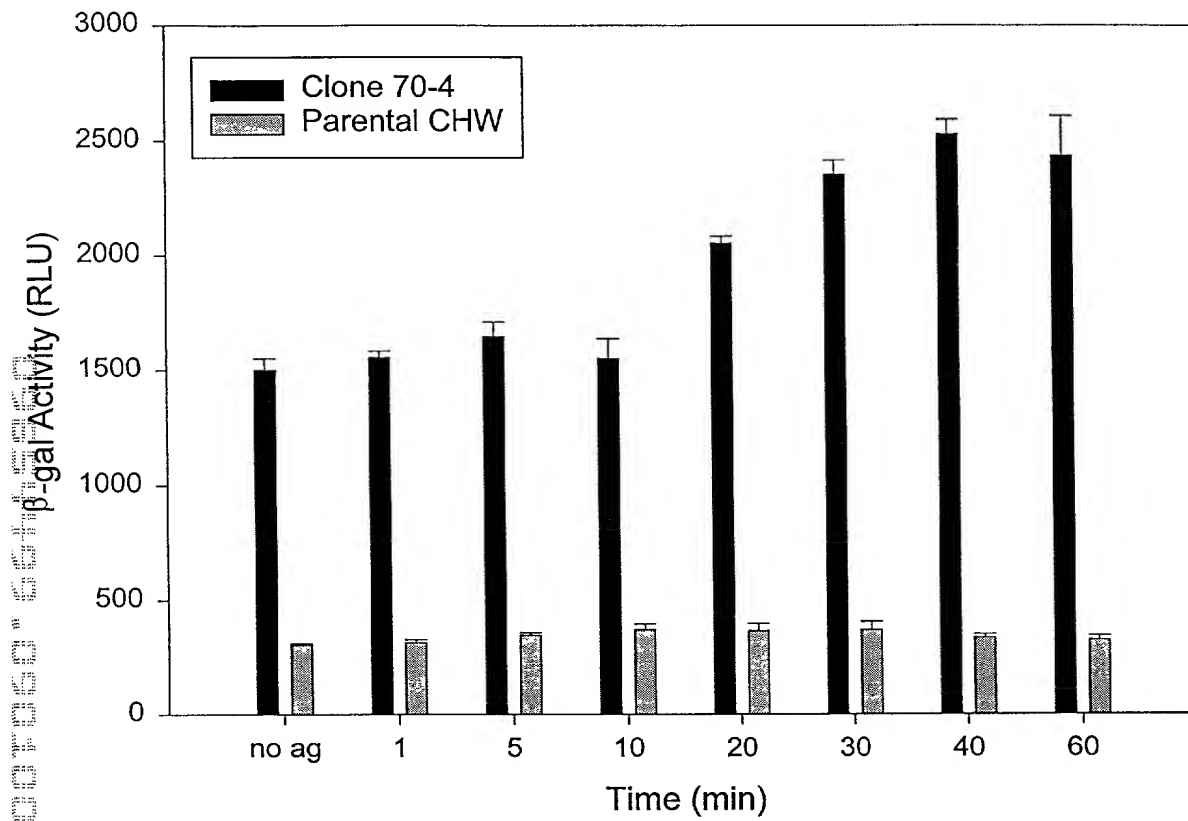


FIGURE 8C

$\beta$ -galactosidase Complementation as a Measurement for  
Adrenergic Receptor Homodimerization in HEK 293 Cells  
Coexpressing  $\beta$ 2AR- $\beta$ gal $\Delta\alpha$  and  $\beta$ 2AR- $\beta$ gal $\Delta\omega$ .

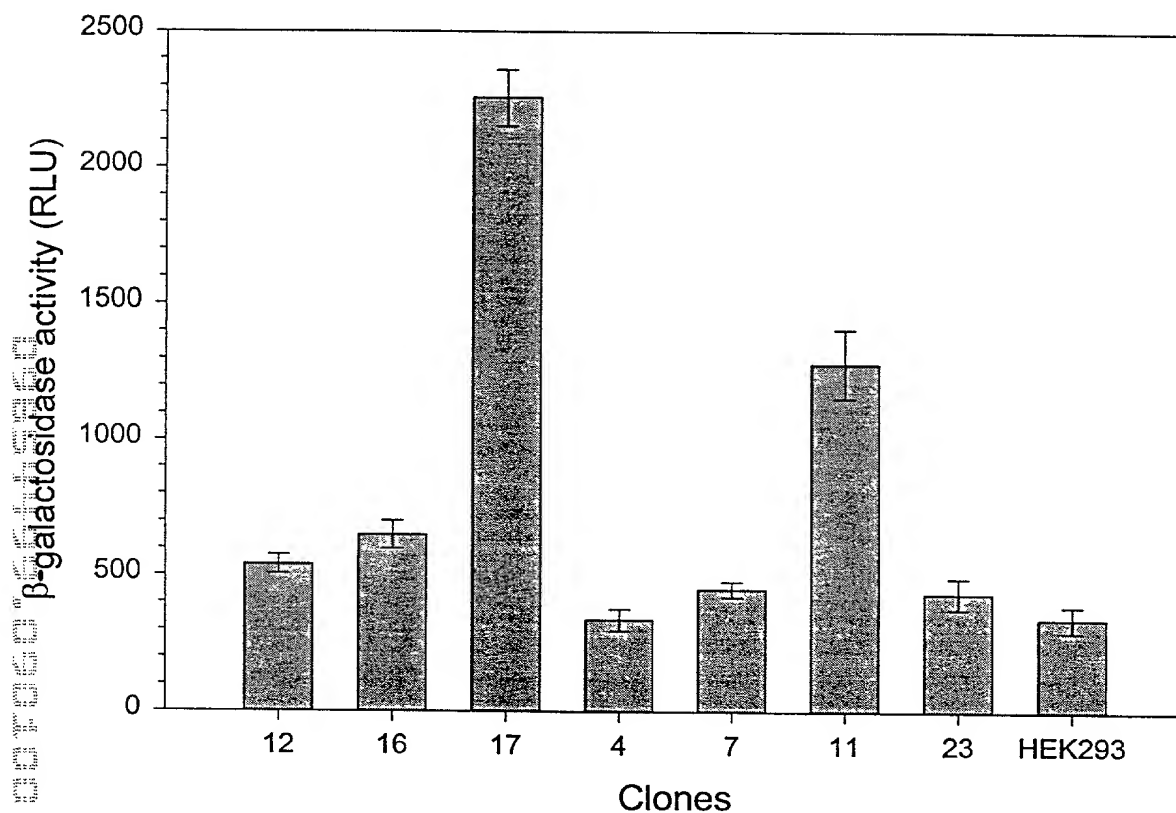


FIGURE 9A

Agonist Stimulated cAMP Response in HEK 293 Cells  
Coexpressing  $\beta 2AR$ - $\beta gal\Delta\alpha$  and  $\beta 2AR$ - $\beta gal\Delta\omega$

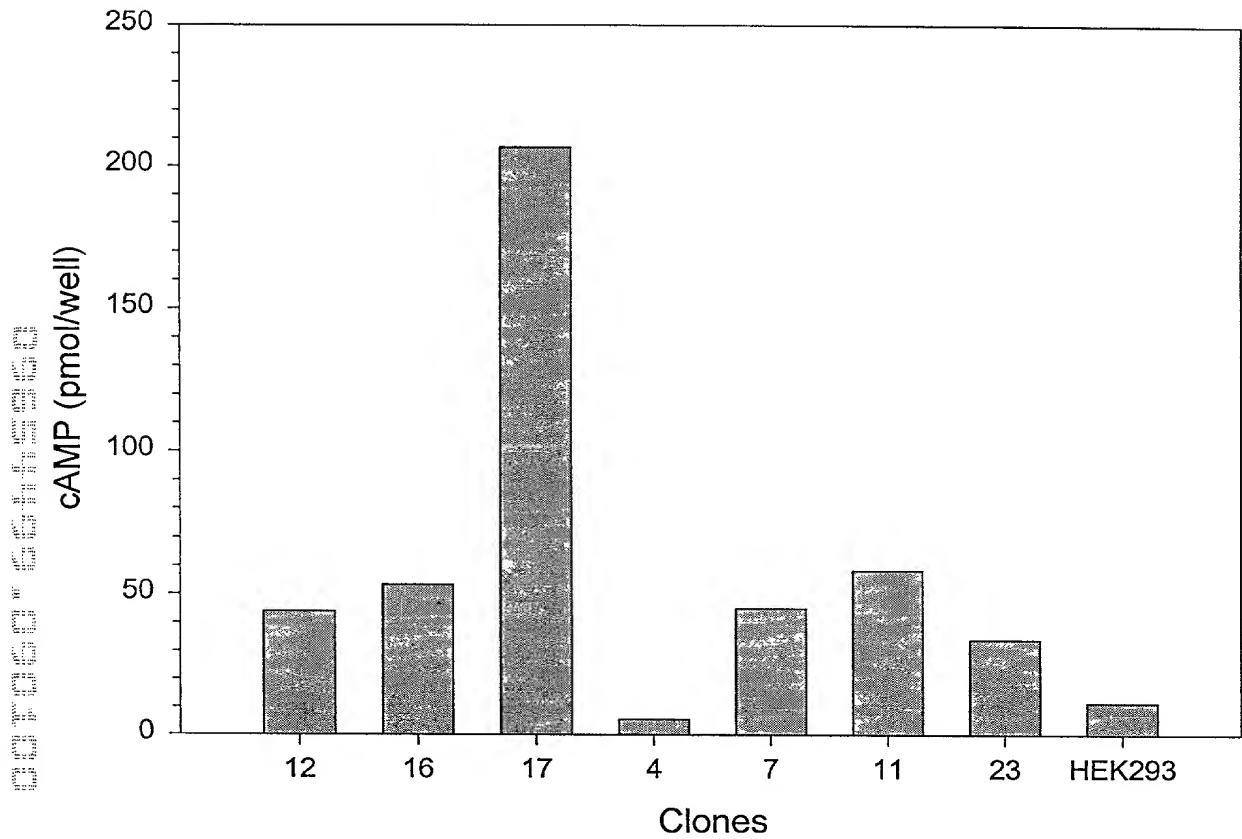


FIGURE 9B

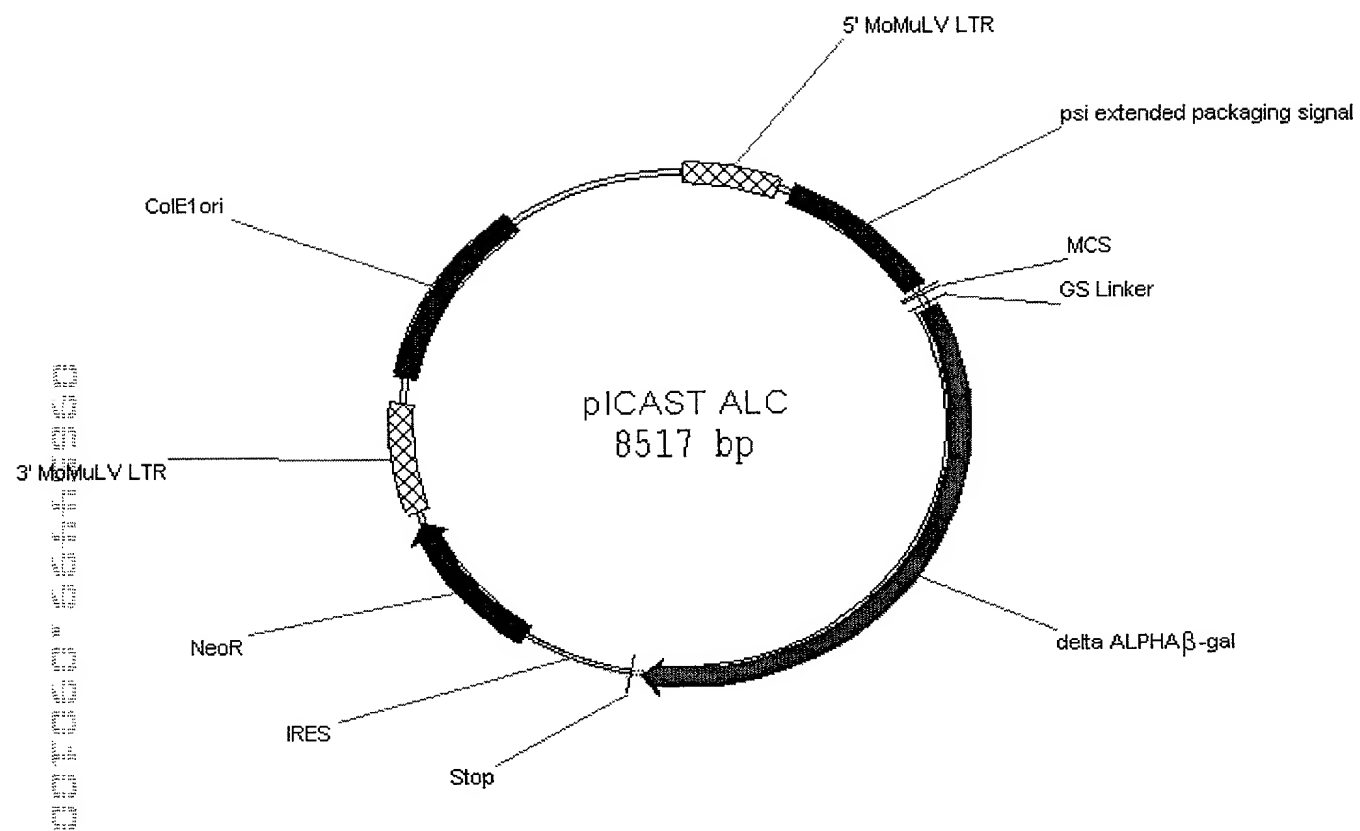


Figure 10A

```

1  CTGCAGCCTG AATATGGGCC AAACAGGATA TCTGTGGTAA GCAGTTCCTG
   GACGTCGGAC TTATACCCCG TTTGTCCTAT AGACACCATT CGTCAAGGAC
-----
51  CCCC GGCTCA GGGCCAAGAA CAGATGGAAC AGCTGAATAT GGGCCAAACA
   GGGGCCGAGT CCCGGTTCTT GTCTACCTTG TCGACTTATA CCCGGTTTGT
-----
101 GGATATCTGT GGTAAGCAGT TCCTGCCCCG GCTCAGGGCC AAGAACAGAT
   CCTATAGACA CCATTTCGTCA AGGACGGGGC CGAGTCCCCG TTCTTGTCTA
-----
151 GGTCCCCAGA TGCGGTCCAG CCCTCAGCAG TTTCTAGAGA ACCATCAGAT
   CCAGGGGTCT ACGCCAGGTC GGGAGTCGTC AAAGATCTCT TGGTAGTCTA
-----
201 GTTTCAGGG TGCCCCAAGG ACCTGAAATG ACCCTGTGCC TTATTTGAAC
   CAAAGGTCCC ACGGGGTTC TGGACTTTAC TGGGACACGG AATAAACTTG
-----
251 TAACCAATCA GTTCGCTTCT CGCTTCTGTT CGCGCGCTTC TGCTCCCCGA
   ATTGGTTAGT CAAGCGAAGA GCGAAGACAA GCGCGCGAAG ACGAGGGGCT
-----
301 GCTCAATAAA AGAGCCCACA ACCCCTCACT CGGGGCGCCA GTCTCCGAT
   CGAGTTATTT TCTCGGTGT TGGGGAGTGA GCGCGCGGT CAGGAGGCTA
-----
351 TGACTGAGTC GCGCGGTAC CCGTGTATCC AATAAACCTT CTTGCAGTTG
   ACTGACTCAG CGGGCCCATG GGCACATAGG TTATTTGGGA GAACGTCAAC
-----
401 CATCCGACTT GTGGTCTCGC TGTTCTTGG GAGGGTCTCC TCTGAGTGAT
   GTAGGCTGAA CACCAGAGCG ACAAGGAACC CTCCCAGAGG AGACTCACTA
-----
451 TGACTACCCG TCAGCGGGGG TCTTTTATTT GGGGGCTCGT CCGGGATCGG
   ACTGATGGGC AGTCGCCCCC AGAAAGTAAA CCCCCGAGCA GGCCCTAGCC
-----
501 GAGACCCCTG CCCAGGGACC ACCGACCCAC CACCGGGAGG CAAGCTGGCC
   CTCTGGGGAC GGGTCCCTGG TGGCTGGGTG GTGGCCCTCC GTTCGACCGG
-----
551 AGCAACTTAT CTGTGTCTGT CCGATTGTCT AGTGTCTATG ACTGATTTTA
   TCGTTGAATA GACACAGACA GGCTAACAGA TCACAGATAC TGACTAAAT
-----
601 TGCGCCTGCG TCGGTACTAG TTAGCTAACT AGCTCTGTAT CTGGCGGACC
   ACGCGGACGC AGCCATGATC AATCGATTGA TCGAGACATA GACCGCTGG
-----
651 CGTGGTGGAA CTGACGAGTT CTGAACACCC GGCCGCAACC CTGGGAGACG
   GCACCACCTT GACTGCTCAA GACTTGTGGG CCGGCGTTGG GACCCTCTGC
-----
701 TCCCAGGGAC TTTGGGGGCC GTTTTGTGG CCCGACCTGA GGAAGGGAGT
   AGGGTCCCCTG AAACCCCCCG CAAAAACACC GGGCTGGACT CCTTCCCTCA
-----
751 CGATGTGGAA TCCGACCCCG TCAGGATATG TGGTTCTGGT AGGAGACGAG
   GCTACACCTT AGGCTGGGGC AGTCCTATAC ACCAAGACCA TCCTCTGCTC
-----
801 AACCTAAAC AGTTCCCGCC TCCGTCTGAA TTTTGTCTT CGGTTTGGAA
   TTGGATTTTG TCAAGGGCGG AGGCAGACTT AAAAACGAAA GCCAAACCTT
-----
851 CCGAAGCCGC GCGTCTTGTC TGCTGCAGCA TCGTTCTGTG TTGTCTCTGT
   GGCTTCGGCG CGCAGAACAG ACGACGTCGT AGCAAGACAC AACAGAGACA
-----
901 CTGACTGTGT TTCTGTATTT GTCTGAAAAT TAGGGCCAGA CTGTTACCAC
   GACTGACACA AAGACATAAA CAGACTTTTA ATCCCGGTCT GACAATGGTG
-----

```

FIGURE 10B

```
951  TCCCTTAAGT TTAGCCTTAG GTAACGGAA AGATGTCGAG CGGCTCGCTC
     AGGGAATTCA AACTGGAATC CATTGACCTT TCTACAGCTC GCCGAGCGAG
-----
1001  ACAACCAGTC GGTAGATGTC AAGAAGAGAC GTTGGGTTAC CTTCTGCTCT
     TGTGCGTCAG CCATCTACAG TTCTTCTCTG CAACCCAATG GAAGACGAGA
-----
1051  GCAGAATGGC CAACCTTTAA CGTCGGATGG CCGCGAGACG GCACCTTTAA
     CGTCTTACCG GTTGGAAATT GCAGCCTACC GCGCCTCTGC CGTGGAAATT
-----
1101  CCGAGACCTC ATCACCCAGG TTAAGATCAA GGTCTTTTCA CCTGGCCCGC
     GGCTCTGGAG TAGTGGGTCC AATTCTAGTT CCAGAAAAGT GGACCGGGCG
-----
1151  ATGGACACCC AGACCAGGTC CCCTACATCG TGACCTGGGA AGCCTTGGCT
     TACCTGTGGG TCTGGTCCAG GGGATGTAGC ACTGGACCCT TCGGAACCGA
-----
1201  TTTGACCCCC CTCCCTGGGT CAAGCCCTTT GTACACCCTA AGCCTCCGCC
     AAACCTGGGG GAGGGACCCA GTTCGGGAAA CATGTGGGAT TCGGAGGCGG
-----
1251  TCCTCTTCCT CCATCCGCCC CGTCTCTCCC CTTTGAACCT CCTCGTTCTG
     AGGAGAAGGA GGTAGGCGGG GCAGAGAGGG GGAACCTGGA GGAGCAAGCT
-----
1301  CCCCgcctcg ATCCTCCCTT TATCCAGCCC TCACTCCTTC TCTAGGCGCC
     GGGGCGGAGC TAGGAGGGAA ATAGGTCGGG AGTGAGGAAG AGATCCGCGG
-----
1351  GGCCGCTCTA GCCCATTAAT ACGACTCACT ATAGGGCGAT TCGAATCAGG
     CCGGCGAGAT CGGGTAATTA TGCTGAGTGA TATCCCGCTA AGCTTAGTCC
-----
1401  CCTTGGCGCG CCGGATCCTT AATTAAGCGC AATTGGGAGG TGGCGGTAGC
     GGAACCGCGC GGCCTAGGAA TTAATTCGCG TTAACCTCC ACCGCCATCG
-----
+2    M G V I T D S L A V V A R T D
     }-----
1451  CTCGAGATGG GCGTGATTAC GGATTCACTG GCCGTCGTGG CCCGCACCGA
     GAGCTCTACC CGCACTAATG CCTAAGTGAC CGGCAGCACC GGGCGTGGCT
-----
+2    R P S Q Q L R S L N G E W R F A
-----
1501  TCGCCCTTCC CAACAGTTAC GCAGCCTGAA TGGCGAATGG CGCTTTGCCT
     AGCGGGAAGG GTTGTCAATG CGTCGGACTT ACCGCTTACC GCGAAACGGA
-----
+2    W F P A P E A V P E S W L E C D L
-----
1551  GGTTTCCGGC ACCAGAAGCG GTGCCGAAA GCTGGCTGGA GTGCGATCTT
     CCAAAGGCCG TGGTCTTCGC CACGGCCTTT CGACCGACCT CACGCTAGAA
-----
+2    P E A D T V V V P S N W Q M H G Y
-----
1601  CCTGAGGCCG ATACTGTCGT CGTCCCCTCA AACTGGCAGA TGCACGGTTA
     GGACTCCGGC TATGACAGCA GCAGGGGAGT TTGACCGTCT ACGTGCCAAT
-----
+2    D A P I Y T N V T Y P I T V N P
-----
1651  CGATGCGCCC ATCTACACCA ACGTGACCTA TCCCATTACG GTCAATCCGC
     GCTACGCGGG TAGATGTGGT TGCACTGGAT AGGGTAATGC CAGTTAGGCG
-----
```

+2 P F V P T E N P T G C Y S L T F N

1701 CGTTTGTTC CACGGAGAAT CCGACGGGTT GTTACTCGCT CACATTTAAT  
GCAAACAAGG GTGCCTCTTA GGCTGCCCAA CAATGAGCGA GTGTAAATTA

+2 V D E S W L Q E G Q T R I I F D G

1751 GTTGATGAAA GCTGGCTACA GGAAGGCCAG ACGCGAATTA TTTTGTATGG  
CAACTACTTT CGACCGATGT CCTTCCGGTC TGCGCTTAAT AAAAATACC

+2 V N S A F H L W C N G R W V G Y

1801 CGTTAACTCG GCGTTTCATC TGTGGTGCAA CGGGCGCTGG GTCGGTTACG  
GCAATTGAGC CGCAAAGTAG ACACCACGTT GCCCGCGACC CAGCCAATGC

+2 G Q D S R L P S E F D L S A F L R

1851 GCCAGGACAG TCGTTTGCCG TCTGAATTTG ACCTGAGCGC ATTTTACGC  
CGGTCTCTGTC AGCAAACGGC AGACTTAAAC TGGACTCGCG TAAAAATGCG

+2 A G E N R L A V M V L R W S D G S

1901 GCCGGAGAAA ACCGCCTCGC GGTGATGGTG CTGCGCTGGA GTGACGGCAG  
CGGCCTCTTT TGGCGGAGCG CCACTACCAC GACGCGACCT CACTGCCGTC

+2 Y L E D Q D M W R M S G I F R D

1951 TTATCTGGAA GATCAGGATA TGTGGCGGAT GAGCGGCATT TTCCGTGACG  
AATAGACCTT CTAGTCCTAT ACACCGCCTA CTCGCGGTAA AAGGCACTGC

+2 V S L L H K P T T Q I S D F H V A

2001 TCTCGTTGCT GCATAAACCG ACTACACAAA TCAGCGATTT CCATGTTGCC  
AGAGCAACGA CGTATTTGGC TGATGTGTTT AGTCGCTAAA GGTACAACGG

+2 T R F N D D F S R A V L E A E V Q

2051 ACTCGCTTTA ATGATGATTT CAGCCGCGCT GTACTGGAGG CTGAAGTTCA  
TGAGCGAAAT TACTACTAAA GTCGGCGCGA CATGACCTCC GACTTCAAGT

+2 M C G E L R D Y L R V T V S L W

2101 GATGTGCGGC GAGTTGCGTG ACTACCTACG GGTAACAGTT TCTTTATGGC  
CTACACGCCG CTCAACGCAC TGATGGATGC CCATTGTCAA AGAAATACCG

+2 Q G E T Q V A S G T A P F G G E I

2151 AGGGTGAAAC GCAGGTGCGC AGCGGCACCG CGCCTTTCGG CGGTGAAATT  
TCCCACTTTG CGTCCAGCGG TCGCCGTGGC GCGGAAAGCC GCCACTTTAA

+2 I D E R G G Y A D R V T L R L N V

2201 ATCGATGAGC GTGGTGGTTA TGCCGATCGC GTCACACTAC GTCTGAACGT  
TAGCTACTCG CACCACCAAT ACGGCTAGCG CAGTGTGATG CAGACTTGCA

+2 E N P K L W S A E I P N L Y R A

2251 CGAAAACCCG AACTGTGGA GCGCCGAAAT CCCGAATCTC TATCGTGCGG  
GCTTTTGGGC TTTGACACCT CGCGGCTTTA GGGCTTAGAG ATAGCACGCC

```
+2 V V E L H T A D G T L I E A E A C
-----
2301 TGGTTGAACT GCACACCGCC GACGGCACGC TGATTGAAGC AGAAGCCTGC
     ACCAACTTGA CGTGTGGCGG CTGCCGTGCG ACTAACTTCG TCTTCGGACG
-----
+2 D V G F R E V R I E N G L L L L N
-----
2351 GATGTCGGTT TCCGCGAGGT GCGGATTGAA AATGGTCTGC TGCTGCTGAA
     CTACAGCCAA AGGCGCTCCA CGCCTAACTT TTACCAGACG ACGACGACTT
-----
+2 G K P L L I R G V N R H E H H P
-----
2401 CGGCAAGCCG TTGCTGATTC GAGGCGTTAA CCGTCACGAG CATCATCCTC
     GCCGTTCCGC AACGACTAAG CTCCGCAATT GGCAGTGCTC GTAGTAGGAG
-----
+2 L H G Q V M D E Q T M V Q D I L L
-----
2451 TGCATGGTCA GGTCAATGGAT GAGCAGACGA TGGTGCAGGA TATCCTGCTG
     ACGTACCACT CCAGTACCTA CTCGTCTGCT ACCACGTCTT ATAGGACGAC
-----
+2 M K Q N N F N A V R C S H Y P N H
-----
2501 ATGAAGCAGA ACAACTTTAA CGCCGTGCGC TGTTTCGATT ATCCGAACCA
     TACTTCGTCT TGTTGAAATT GCGGCACGCG ACAAGCGTAA TAGGCTTGGT
-----
+2 P L W Y T L C D R Y G L Y V V D
-----
2551 TCCGCTGTGG TACACGCTGT GCGACCGCTA CGGCCTGTAT GTGGTGGATG
     AGGCGACACC ATGTGCGACA CGCTGGCGAT GCCGGACATA CACCACCTAC
-----
+2 E A N I E T H G M V P M N R L T D
-----
2601 AAGCCAATAT TGAAACCCAC GGCATGGTGC CAATGAATCG TCTGACCGAT
     TTCGGTTATA ACTTTGGGTG CCGTACCACG GTTACTTAGC AGACTGGCTA
-----
+2 D P R W L P A M S E R V T R M V Q
-----
2651 GATCCGCGCT GGCTACCGGC GATGAGCGAA CGCGTAACGC GAATGGTGCA
     CTAGGCGCGA CCGATGGCCG CTACTCGCTT GCGCATTGCG CTTACCACGT
-----
+2 R D R N H P S V I I W S L G N E
-----
2701 GCGCGATCGT AATCACCCGA GTGTGATCAT CTGGTCGCTG GGAATGAAT
     CGCGCTAGCA TTAGTGGGCT CACACTAGTA GACCAGCGAC CCCTTACTTA
-----
+2 S G H G A N H D A L Y R W I K S V
-----
2751 CAGGCCACGG CGCTAATCAC GACGCGCTGT ATCGCTGGAT CAAATCTGTC
     GTCCGGTGCC GCGATTAGTG CTGCGCGACA TAGCGACCTA GTTTAGACAG
-----
+2 D P S R P V Q Y E G G G A D T T A
-----
2801 GATCCTTCCC GCCCGGTGCA GTATGAAGGC GCGGAGCCG ACACCACGGC
     CTAGGAAGGG CGGGCCACGT CATACTTCCG CCGCCTCGGC TGTGGTGCCG
-----
+2 T D I I C P M Y A R V D E D Q P
-----
2851 CACCGATATT ATTTGCCCGA GTACGCGCG CGTGGATGAA GACCAGCCCT
     GTGGCTATAA TAAACGGGCT ACATGCGCGC GCACCTACTT CTGGTCGGGA
```



+2 F P A V P K W S I K K W L S L P G

2901 TCCCGGCTGT GCCGAAATGG TCCATCAAAA AATGGCTTTC GCTACCTGGA  
AGGGCCGACA CGGCTTTACC AGGTAGTTTT TTACCGAAAG CGATGGACCT

+2 E T R P L I L C E Y A H A M G N S

2951 GAGACGCGCC CGCTGATCCT TTGCGAATAC GCCCAGCGCA TGGGTAACAG  
CTCTGCGCGG GCGACTAGGA AACGCTTATG CGGGTGCGCT ACCCATTGTC

+2 L G G F A K Y W Q A F R Q Y P R

3001 TCTTGGCGGT TTCGCTAAAT ACTGGCAGGC GTTTCGTCAG TATCCCCGTT  
AGAACCGCCA AAGCGATTTA TGACCGTCCG CAAAGCAGTC ATAGGGGCAA

+2 L Q G G F V W D W V D Q S L I K Y

3051 TACAGGGCGG CTTCGTCTGG GACTGGGTGG ATCAGTCGCT GATTAAATAT  
ATGTCCCGCC GAAGCAGACC CTGACCCACC TAGTCAGCGA CTAATTTATA

+2 D E N G N P W S A Y G G D F G D T

3101 GATGAAAACG GCAACCCGTG GTCGGCTTAC GGCGGTGATT TTGGCGATAC  
CTACTTTTGC CGTTGGGCAC CAGCCGAATG CCGCCACTAA AACCGCTATG

+2 P N D R Q F C M N G L V F A D R

3151 GCCGAACGAT CGCCAGTTCT GTATGAACGG TCTGGTCTTT GCCGACCGCA  
CGGCTTGCTA GCGGTCAAGA CATACTTGCC AGACCAGAAA CGGCTGGCGT

+2 T P H P A L T E A K H Q Q Q F F Q

3201 CGCCGCATCC AGCGCTGACG GAAGCAAAAC ACCAGCAGCA GTTTTTCCAG  
GCGGCGTAGG TCGCGACTGC CTTCGTTTTG TGGTCGTCGT CAAAAAGGTC

+2 F R L S G Q T I E V T S E Y L F R

3251 TTCCGTTTAT CCGGGCAAAC CATCGAAGTG ACCAGCGAAT ACCTGTTCCG  
AAGGCAAATA GGCCCGTTTG GTAGCTTCAC TGGTCGCTTA TGGACAAGGC

+2 H S D N E L L H W M V A L D G K

3301 TCATAGCGAT AACGAGCTCC TGCCTGGAT GGTGGCGCTG GATGGTAAGC  
AGTATCGCTA TTGCTCGAGG ACGTGACCTA CCACCGCGAC CTACCATTCTG

+2 P L A S G E V P L D V A P Q G K Q

3351 CGCTGGCAAG CGGTGAAGTG CCTCTGGATG TCGCTCCACA AGGTAAACAG  
GCGACCGTTC GCCACTTCAC GGAGACCTAC AGCGAGGTGT TCCATTTGTC

+2 L I E L P E L P Q P E S A G Q L W

3401 TTGATTGAAC TGCCTGAACT ACCGCAGCCG GAGAGCGCCG GGCAACTCTG  
AACTAACTTG ACGGACTTGA TGGCGTCGGC CTCTCGCGGC CCGTTGAGAC

+2 L T V R V V Q P N A T A W S E A

3451 GCTCACAGTA CGCGTAGTGC AACCGAACGC GACCGCATGG TCAGAAGCCG  
CGAGTGTAT GCGCATCAG TTGGCTTGCG CTGGCGTACC AGTCTTCGGC

```
+2 G H I S A W Q Q W R L A E N L S V
-----
3501 GGCACATCAG CGCCTGGCAG CAGTGGCGTC TGGCGGAAAA CTCAGTGTG
    CCGTG TAGTC GCGGACCGTC GTCACCGCAG ACCGCCTTTT GGAGTCACAC
-----
+2 T L P A A S H A I P H L T T S E M
-----
3551 ACGCTCCCCG CCGCGTCCCA CGCCATCCCG CATCTGACCA CCAGCGAAAT
    TGCGAGGGGC GGCGCAGGGT GCGGTAGGGC GTAGACTGGT GGTGCGCTTA
-----
+2 D F C I E L G N K R W Q F N R Q
-----
3601 GGATTTTTGC ATCGAGCTGG GTAATAAGCG TTGGCAATTT AACCGCCAGT
    CCTAAAAACG TAGCTCGACC CATTATTCGC AACCGTTAAA TTGGCGGTCA
-----
+2 S G F L S Q M W I G D K K Q L L T
-----
3651 CAGGCTTTCT TTCACAGATG TGGATTGGCG ATAAAAACA ACTGCTGACG
    GTCCGAAAGA AAGTGTCTAC ACCTAACCGC TATTTTTTGT TGACGACTGC
-----
+2 P L R D Q F T R A P L D N D I G V
-----
3701 CCGCTGCGCG ATCAGTTCAC CCGTGACCG CTGGATAACG ACATTGGCGT
    GCGCAGCGCG TAGTCAAGTG GGCACGTGGC GACCTATTGC TGTAAACGCA
-----
+2 S E A T R I D P N A W V E R W K
-----
3751 AAGTGAAGCG ACCCGCATTG ACCCTAACGC CTGGGTCGAA CGCTGGAAGG
    TTCATTTCGC TGGGCGTAAC TGGGATTGCG GACCCAGCTT GCGACCTTCC
-----
+2 A A G H Y Q A E A A L L Q C T A D
-----
3801 CGGCGGGCCA TTACCAGGCC GAAGCAGCGT TGTTGCAGTG CACGGCAGAT
    GCCGCCCGGT AATGGTCCGG CTTCTGTCGCA ACAACGTCAC GTGCCGTCTA
-----
+2 T L A D A V L I T T A H A W Q H Q
-----
3851 ACACTTGCTG ATGCGGTGCT GATTACGACC GCTCACGCGT GGCAGCATCA
    TGTGAACGAC TACGCCACGA CTAATGCTGG CGAGTGC GCA CCGTCGTAGT
-----
+2 G K T L F I S R K T Y R I D G S
-----
3901 GGGGAAAACC TTATTATCA GCCGAAAAC CTACCGGATT GATGGTAGTG
    CCCCTTTTGG AATAAATAGT CGGCCTTTTG GATGGCCTAA CTACCATCAC
-----
+2 G Q M A I T V D V E V A S D T P H
-----
3951 GTCAAATGGC GATTACCGTT GATGTTGAAG TGGCGAGCGA TACACCGCAT
    CAGTTTACCG CTAATGGCAA CTACAAC TTC ACCGCTCGCT ATGTGGCGTA
-----
+2 P A R I G L N C Q L A Q V A E R V
-----
4001 CCGGCGCGGA TTGGCCTGAA CTGCCAGCTG GCGCAGGTAG CAGAGCGGGT
    GGCCGCGCCT AACCGGACTT GACGGTCGAC CGCTCCATC GTCTCGCCCA
-----
+2 N W L G L G P Q E N Y P D R L T
-----
4051 AAATGGCTC GGATTAGGGC CGCAAGAAAA CTATCCCGAC CGCCTTACTG
    TTTGACCGAG CTAATCCCG GCGTTCTTTT GATAGGGCTG GCGGAATGAC
-----
```

```
+2 A A C F D R W D L P L S D M Y T P
-----
4101 CCGCCTGTTT TGACCGCTGG GATCTGCCAT TGTCAGACAT GTATACCCCG
      GCGGACAAAA ACTGGCGACC CTAGACGGTA ACAGTCTGTA CATATGGGGC
-----
+2 Y V F P S E N G L R C G T R E L N
-----
4151 TACGTCTTCC CGAGCGAAAA CGGTCTGCGC TCGGGGACGC GCGAATTGAA
      ATGCAGAAGG GCTCGCTTTT GCCAGACGCG ACGCCCTGCG CGCTTAACTT
-----
+2 Y G P H Q W R G D F Q F N I S R
-----
4201 TTATGGCCCA CACCAGTGGC GCGGCGACTT CCAGTTCAAC ATCAGCCGCT
      AATACCGGGT GTGGTCACCG CGCCGCTGAA GGTCAAGTTG TAGTCGGCGA
-----
+2 Y S Q Q Q L M E T S H R H L L H A
-----
4251 ACAGTCAACA GCAACTGATG GAAACCAGCC ATCGCCATCT GCTGCACGCG
      TGTCAGTTGT CGTTGACTAC CTTTGGTCCG TAGCGGTAGA CGACGTGCGC
-----
+2 E E G T W L N I D G F H M G I G G
-----
4301 GAAGAAGGCA CATGGCTGAA TATCGACGGT TTCCATATGG GGATTGGTGG
      CTTCTTCCGT GTACCGACTT ATAGCTGCCA AAGGTATAACC CCTAACCACC
-----
+2 D D S W S P S V S A E F Q L S A
-----
4351 CGACGACTCC TGGAGCCCGT CAGTATCGGC GGAATTCCAG CTGAGCGCCG
      GCTGCTGAGG ACCTCGGGCA GTCATAGCCG CCTTAAGGTC GACTCGCGGC
-----
+2 G R Y H Y Q L V W C Q K R S D Y K
-----
4401 GTCGCTACCA TTACCAAGTG GTCTGGTGTG AAAAAAGATC TGACTATAAA
      CAGCGATGGT AATGGTCAAC CAGACCACAG TTTTCTCTAG ACTGATATTT
-----
+2 D E D L D H H H H H H R
-----
4451 GATGAGGACC TCGACCATCA TCATCATCAT CACCGGTAAT AATAGGTAGA
      CTACTCCTGG AGCTGGTAGT AGTAGTAGTA GTGGCCATTA TTATCCATCT
-----
+2
-----
4501 TAAGTGACTG ATTAGATGCA TTGATCCCTC GACCAATTCC GGTATTATTTT
      ATTCACTGAC TAATCTACGT AACTAGGGAG CTGGTTAAGG CCAATAAAAG
-----
4551 CACCATATTG CCGTCTTTTG GCAATGTGAG GGCCCGGAAA CCTGGCCCTG
      GTGGTATAAC GGCAGAAAAC CGTTACACTC CCGGGCCTTT GGACCGGGAC
-----
4601 TCTTCTTGAC GAGCATTCCT AGGGGTCTTT CCCCTCTCGC CAAAGGAATG
      AGAAGAACTG CTCGTAAGGA TCCCCAGAAA GGGGAGAGCG GTTTCCTTAC
-----
4651 CAAGGTCTGT TGAATGTCGT GAAGGAAGCA GTTCCTCTGG AAGCTTCTTG
      GTTCCAGACA ACTTACAGCA CTTCTTCTGT CAAGGAGACC TTCGAAGAAC
-----
4701 AAGACAAACA ACGTCTGTAG CGACCCTTTG CAGGCAGCGG AACCCCCAC
      TTCTGTTTGT TGCAGACATC GCTGGGAAAC GTCCGTCTGCC TTGGGGGGTG
-----
4751 CTGGCGACAG GTGCCTCTGC GGCCAAAAGC CACGTGTATA AGATACACCT
      GACCGCTGTC CACGGAGACG CCGGTTTTCG GTGCACATAT TCTATGTGGA
-----
```

4801 GCAAAGGCGG CACAACCCCA GTGCCACGTT GTGAGTTGGA TAGTTGTGGA  
CGTTTCCGCC GTGTTGGGGT CACGGTGCAA CACTCAACCT ATCAACACCT

-----

4851 AAGAGTCAAA TGGCTCTCCT CAAGCGTATT CAACAAGGGG CTGAAGGATG  
TTCTCAGTTT ACCGAGAGGA GTTCGCATAA GTTGTTCCCC GACTTCCTAC

-----

4901 CCCAGAAGGT ACCCCATTGT ATGGGATCTG ATCTGGGGCC TCGGTGCACA  
GGGTCTTCCA TGGGGTAACA TACCCTAGAC TAGACCCCGG AGCCACGTGT

-----

4951 TGCTTTACAT GTGTTTAGTC GAGGTTAAAA AACGTCTAGG CCCCCGAAC  
ACGAAATGTA CACAAATCAG CTCCAATTTT TTGCAGATCC GGGGGGCTTG

-----

5001 CACGGGGACG TGGTTTTCCT TTGAAAAACA CGATGATAAT ACCATGATTG  
GTGCCCCCTGC ACCAAAAGGA AACTTTTGT GCTACTATTA TGGTACTAAC

-----

5051 AACAAAGATGG ATTGCACGCA GGTTCCTCCG CCGCTTGGGT GGAGAGGCTA  
TTGTTCTACC TAACGTGCGT CCAAGAGGCC GCGAACCCTA CCTCTCCGAT

-----

5101 TTCGGCTATG ACTGGGCACA ACAGACAATC GGCTGCTCTG ATGCCGCCGT  
AAGCCGATAC TGACCCGTGT TGTCTGTTAG CCGACGAGAC TACGGCGGCA

-----

5151 GTTCGGCTG TCAGCGCAGG GCGCGCCGGT TCTTTTGTG AAGACCGACC  
CAAGCCGAC AGTCGCGTCC CCGCGGGCCA AGAAAAACAG TTCTGGCTGG

-----

5201 TGTCCGGTGC CCTGAATGAA CTGCAGGACG AGGCAGCGCG GCTATCGTGG  
ACAGGCCACG GGACTTACTT GACGTCTGTC TCCGTCGCGC CGATAGCACC

-----

5251 CTGGCCACGA CGGGCGTTCC TTGCGCAGCT GTGCTCGACG TTGTCACTGA  
GACCGGTGCT GCGCGCAAGG AACGCGTCGA CACGAGCTGC AACAGTGACT

-----

5301 AGCGGGAAGG GACTGGCTGC TATTGGGCGA AGTGCCGGGG CAGGATCTCC  
TCGCCCTTCC CTGACCGACG ATAACCCGCT TCACGGCCCC GTCCTAGAGG

-----

5351 TGTCATCTCA CCTTGCTCCT GCCGAGAAAG TATCCATCAT GGCTGATGCA  
ACAGTAGAGT GGAACGAGGA CGGCTCTTTC ATAGGTAGTA CCGACTACGT

-----

5401 ATGCGGCGGC TGCATACGCT TGATCCGGCT ACCTGCCCAT TCGACCACCA  
TACGCCGCCG ACGTATGCGA ACTAGGCCGA TGGACGGGTA AGCTGGTGTT

-----

5451 AGCGAAACAT CGCATCGAGC GAGCACGTAC TCGGATGGAA GCCGGTCTTG  
TCGCTTTGTA GCGTAGCTCG CTCGTGCATG AGCCTACCTT CGGCCAGAAC

-----

5501 TCGATCAGGA TGATCTGGAC GAAGAGCATC AGGGGCTCGC GCCAGCCGAA  
AGCTAGTCCT ACTAGACCTG CTTCTCGTAG TCCCCGAGCG CGGTCGGCTT

-----

5551 CTGTTCGCCA GGCTCAAGGC GCGCATGCCC GACGCGGAGG ATCTCGTCGT  
GACAAGCGGT CCGAGTTCCG CGCGTACGGG CTGCCGCTCC TAGAGCAGCA

-----

5601 GACCCATGGC GATGCCTGCT TGCCGAATAT CATGGTGGAA AATGGCCGCT  
CTGGGTACCG CTACGGACGA ACGGCTTATA GTACCACCTT TTACGGCGA

-----

5651 TTTCTGGATT CATCGACTGT GGCCGGCTGG GTGTGGCGGA CCGCTATCAG  
AAAGACCTAA GTAGCTGACA CCGGCCGACC CACACCGCCT GGCGATAGTC

-----

5701 GACATAGCGT TGGCTACCCG TGATATTGCT GAAGAGCTTG GCGGCGAATG  
CTGTATCGCA ACCGATGGGC ACTATAACGA CTTCTCGAAC CGCCGCTTAC

-----

5751 GGCTGACCGC TTCCTCGTGC TTTACGGTAT CGCCGCTCCC GATTTCGCAGC  
CCGACTGGCG AAGGAGCACG AAATGCCATA GCGGCGAGGG CTAAGCGTCG

5801 GCATCGCCTT CTATCGCCTT CTTGACGAGT TCTTCTGAGC GGGACTCTGG  
CGTAGCGGAA GATAGCGGAA GAACTGCTCA AGAAGACTCG CCCTGAGACC

5851 GGTTCGCATC GATAAAATAA AAGATTTTAT TTAGTCTCCA GAAAAAGGGG  
CCAAGCGTAG CTATTTTATT TTCTAAAATA AATCAGAGGT CTTTTTCCCC

5901 GGAATGAAAG ACCCCACCTG TAGGTTTGGC AAGCTAGCTT AAGTAACGCC  
CCTTACTTTC TGGGTTGGAC ATCCAAACCG TTCGATCGAA TTCATTGCGG

5951 ATTTTGCAAG GCATGGAAAA ATACATAACT GAGAATAGAG AAGTTCAGAT  
TAAAACGTTT CGTACCTTTT TATGTATTGA CTCTTATCTC TTCAAGTCTA

6001 CAAGGTCAGG AACAGATGGA ACAGCTGAAT ATGGGCCAAA CAGGATATCT  
GTTCCAGTCC TTGTCTACCT TGTCGACTTA TACCCGGTTT GTCCTATAGA

6051 GTGTAAGCA GTTCCTGCCC CGGCTCAGGG CCAAGAACAG ATGGAACAGC  
CACCATTTCG CAAGGACGGG GCCGAGTCCC GTTCTTGTC TACCTTGTCG

6101 TGAATATGGG CCAAACAGGA TATCTGTGGT AAGCAGTTCC TGCCCCGGCT  
ACTTATACCC GGTTCGCTCT ATAGACACCA TTCGTCAAGG ACGGGGCCGA

6151 CAGGGCCAAG AACAGATGGT CCCCAGATGC GGTCCAGCCC TCAGCAGTTT  
GTCCCGGTTT TTGTCTACCA GGGGTCTACG CCAGGTCGGG AGTCGTCAA

6201 CTAGAGAACC ATCAGATGTT TCCAGGGTGC CCCAAGGACC TGAAATGACC  
GATCTCTTGG TAGTCTACAA AGGTCCCACG GGGTTCCTGG ACTTTACTGG

6251 CTGTGCCTTA TTTGAACTAA CCAATCAGTT CGCTTCTCGC TTCTGTTTCG  
GACACGGAAT AAACCTGATT GGTAGTCAA GCGAAGAGCG AAGACAAGCG

6301 GCGCTTCTGC TCCCCGAGCT CAATAAAAGA GCCACAACC CCTCACTCGG  
CGCGAAGACG AGGGGCTCGA GTTATTTTCT CGGGTGTGG GGAGTGAGCC

6351 GCGGCCAGTC CTCCGATTGA CTGAGTCGCC CGGGTACCCG TGTATCCAAT  
CCGCGGTCAG GAGGCTAACT GACTCAGCGG GCCCATGGGC ACATAGGTTA

6401 AAACCCTCTT GCAGTTGCAT CCGACTTGTG GTCTCGCTGT TCCTTGGGAG  
TTTGGGAGAA CGTCAACGTA GGCTGAACAC CAGAGCGACA AGGAACCCTC

6451 GGTCTCCTCT GAGTGATTGA CTACCCGTCA GCGGGGGTCT TTCATTATG  
CCAGAGGAGA CTCACTAACT GATGGGCAGT CGCCCCAGA AAGTAAGTAC

6501 CAGCATGTAT CAAAATTAAT TTGGTTTTTT TTCTTAAGTA TTTACATTAA  
GTCGTACATA GTTTTAATTA AACCAAAAAA AAGAATTCAT AAATGTAATT

6551 ATGGCCATAG TTGCATTAAT GAATCGGCCA ACGCGCGGGG AGAGGCGGTT  
TACCGGTATC AACGTAATTA CTTAGCCGGT TCGCGGCCCC TCTCCGCCAA

6601 TCGTATTGG CGCTCTCCG CTTCTCGCT CACTGACTCG CTGCGCTCGG  
ACGCATAACC GCGAGAAGGC GAAGGAGCGA GTGACTGAGC GACGCGAGCC

6651 TCGTTCGGCT GCGGCGAGCG GTATCAGCTC ACTCAAAGGC GGTAATACGG  
AGCAAGCCGA CGCCGCTCGC CATAGTCGAG TGAGTTTCCG CCATTATGCC

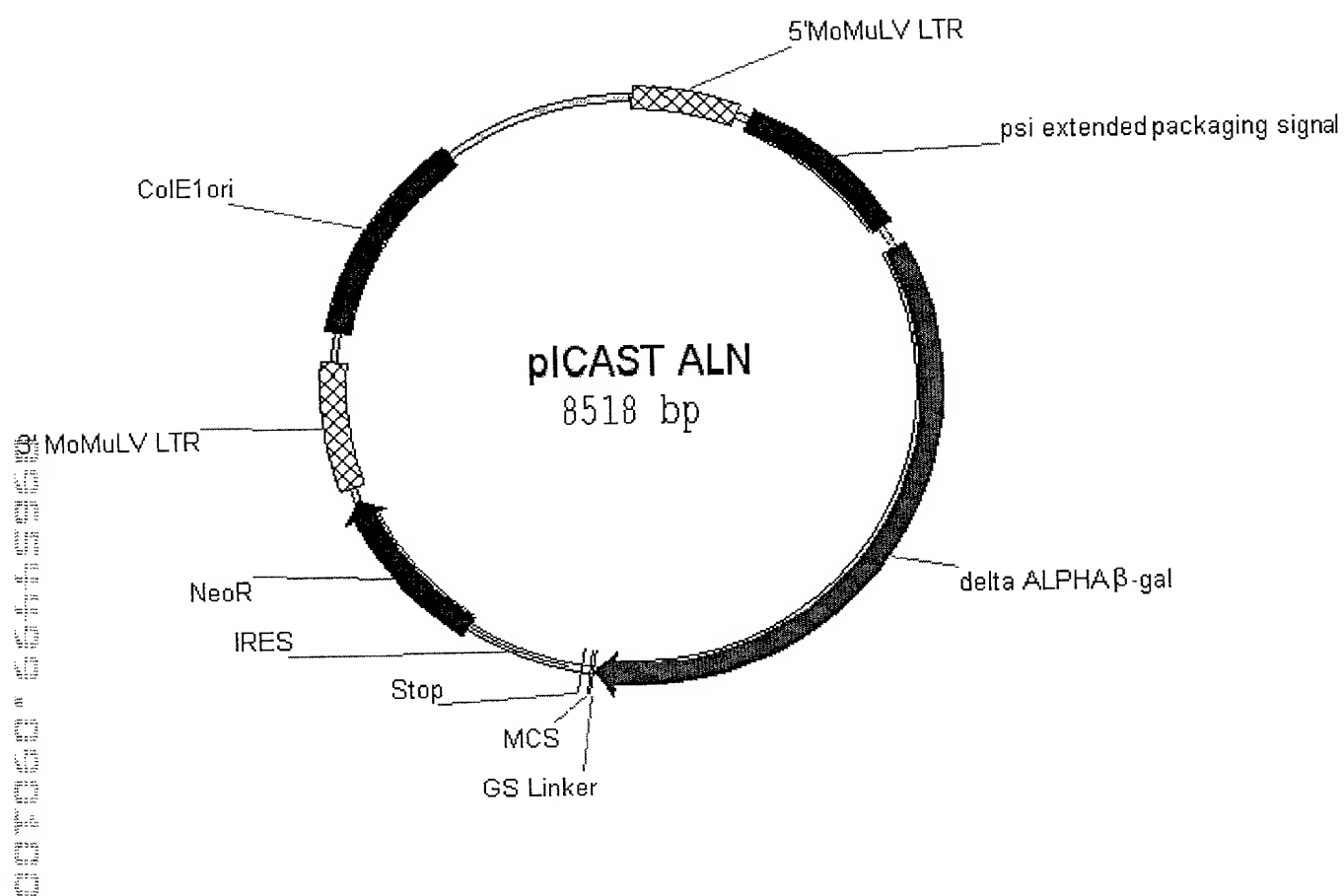


Figure 11A

```

1  CTGCAGCCTG AATATGGGCC AAACAGGATA TCTGTGGTAA GCAGTTCCTG
   GACGTCGGAC TTATACCCGG TTTGTCCTAT AGACACCATT CGTCAAGGAC
-----
51  CCCCCGGCTCA GGGCCAAGAA CAGATGGAAC AGCTGAATAT GGGCCAAACA
   GGGGCCGAGT CCCGGTTCTT GTCTACCTTG TCGACTTATA CCCGGTTTGT
-----
101 GGATATCTGT GGTAAGCAGT TCCTGCCCCG GCTCAGGGCC AAGAACAGAT
   CCTATAGACA CCATTTCGTC AGGACGGGGC CGAGTCCCGG TTCTTGTCTA
-----
151 GGTCCCCAGA TGCGGTCCAG CCCTCAGCAG TTTCTAGAGA ACCATCAGAT
   CCAGGGGTCT ACGCCAGGTC GGGAGTCGTC AAAGATCTCT TGGTAGTCTA
-----
201 GTTTCAGGG TGCCCCAAGG ACCTGAAATG ACCCTGTGCC TTATTTGAAC
   CAAAGGTCCC ACGGGGTTC TGGACTTTAC TGGGACACGG AATAAACTTG
-----
251 TAACCAATCA GTTCGCTTCT CGCTTCTGTT CGCGCGCTTC TGCTCCCCGA
   ATTGGTTAGT CAAGCGAAGA GCGAAGACAA GCGCGCGAAG ACGAGGGGCT
-----
301 GCTCAATAAA AGAGCCCACA ACCCCTCACT CGGGGCGCCA GTCCTCCGAT
   CGAGTTATTT TCTCGGGTGT TGGGGAGTGA GCCCCGCGGT CAGGAGGCTA
-----
351 TGACTGAGTC GCGCGGTAC CCGTGTATCC AATAAACCTT CTGTCAGTTG
   ACTGACTCAG CGGGCCCATG GGCACATAGG TTATTTGGGA GAACGTCAAC
-----
401 CATCCGACTT GTGGTCTCGC TGTTCTTGG GAGGGTCTCC TCTGAGTGAT
   GTAGGCTGAA CACCAGAGCG ACAAGGAACC CTCCCAGAGG AGACTCACTA
-----
451 TGACTACCCG TCAGCGGGGG TCTTTCATTT GGGGGCTCGT CCGGGATCGG
   ACTGATGGGC AGTCGCCCCC AGAAAGTAA CCCCCGAGCA GGCCCTAGCC
-----
501 GAGACCCCTG CCCAGGGACC ACCGACCCAC CACCGGGAGG CAAGCTGGCC
   CTCTGGGGAC GGGTCCCTGG TGGCTGGGTG GTGGCCCTCC GTTCGACCGG
-----
551 AGCAACTTAT CTGTGTCTGT CCGATTGTCT AGTGTCTATG ACTGATTTTA
   TCGTTGAATA GACACAGACA GGCTAACAGA TCACAGATAC TGAATAAAAT
-----
601 TGCGCCTGCG TCGGTACTAG TTAGCTAACT AGCTCTGTAT CTGGCGGACC
   ACGCGGACGC AGCCATGATC AATCGATTGA TCGAGACATA GACCGCCTGG
-----
651 CGTGGTGGAA CTGACGAGTT CTGAACACCC GGCCGCAACC CTGGGAGACG
   GCACCACCTT GACTGCTCAA GACTTGTGGG CCGGCGTTGG GACCCTCTGC
-----
701 TCCAGGGGAC TTTGGGGGCC GTTTTGTGG CCCGACCTGA GGAAGGGAGT
   AGGGTCCCTG AAACCCCGG CAAAACACC GGGCTGGACT CCTTCCCTCA
-----
751 CGATGTGGAA TCCGACCCCG TCAGGATATG TGGTTCTGGT AGGAGACGAG
   GCTACACCTT AGGCTGGGGC AGTCCTATAC ACCAAGACCA TCCTCTGCTC
-----
801 AACCTAAAAC AGTTCCCGCC TCCGTCTGAA TTTTGTGCTT CGGTTTGGAA
   TTGGATTTTG TCAAGGGCGG AGGCAGACTT AAAACGAAA GCCAAACCTT
-----
851 CCGAAGCCGC GCGTCTTGTG TGCTGCAGCA TCGTCTGTG TTGTCTCTGT
   GGCTTCGGCG CGCAGAACAG ACGACGTCGT AGCAAGACAC AACAGAGACA
-----
901 CTGACTGTGT TTCTGTATTT GTCTGAAAAT TAGGGCCAGA CTGTTACCAC
   GACTGACACA AAGACATAAA CAGACTTTTA ATCCCGGTCT GACAATGGTG
-----

```

FIGURE 11B

```
951  TCCCTTAAGT TTGACCTTAG GTAACGGAA AGATGTCGAG CGGCTCGCTC
     AGGGAATTCA AACTGGAATC CATTGACCTT TCTACAGCTC GCCGAGCGAG
-----
1001  ACAACCAGTC GGTAGATGTC AAGAAGAGAC GTTGGGTTAC CTTCTGCTCT
     TGTGGTTCAG CCATCTACAG TTCTTCTCTG CAACCCAATG GAAGACGAGA
-----
1051  GCAGAATGGC CAACCTTTAA CGTCGGATGG CCGCGAGACG GCACCTTTAA
     CGTCTTACCG GTTGGAATTT GCAGCCTACC GGCGCTCTGC CGTGGAATTT
-----
1101  CCGAGACCTC ATCACCAGG TTAAGATCAA GGTCTTTTCA CCTGGCCCGC
     GGCTCTGGAG TAGTGGGTCC AATTCTAGTT CCAGAAAAGT GGACCGGGCG
-----
1151  ATGGACACCC AGACCAGGTC CCCTACATCG TGACCTGGGA AGCCTTGGCT
     TACCTGTGGG TCTGGTCCAG GGGATGTAGC ACTGGACCCT TCGGAACCGA
-----
1201  TTTGACCCCC CTCCCTGGGT CAAGCCCTTT GTACACCCTA AGCCTCCGCC
     AAACCTGGGGG GAGGGACCCA GTTCGGGAAA CATGTGGGAT TCGGAGGCGG
-----
1251  TCCTCTTCCT CCATCCGCCC CGTCTCTCCC CCTTGAACCT CCTCGTTCTG
     AGGAGAAGGA GGTAGGCGGG GCAGAGAGGG GGAACCTTGA GGAGCAAGCT
-----
1301  CCGCGCTCTG ATCCTCCCTT TATCCAGCCC TCACTCCTTC TCTAGGCGCC
     GGGGCGGAGC TAGGAGGGAA ATAGGTCGGG AGTGAGGAAG AGATCCGCGG
-----
1351  GGCCGCTCTA GCCCATTAAT ACGACTCACT ATAGGGCGAT TCGAACACCA
     CCGGCGAGAT CGGGTAATTA TGCTGAGTGA TATCCCGCTA AGCTTGTGGT
-----
1401  TGCACCATCA TCATCATCAC GTCGACTATA AAGATGAGGA CCTCGAGATG
     ACGTGGTAGT AGTAGTAGTG CAGCTGATAT TTCTACTCCT GGAGCTCTAC
-----
1451  GGCGTGATTA CGGATTCACG GGCCGTCGTG GCGCGCACCG ATCGCCCTTC
     CCGCACTAAT GCCTAAGTGA CCGGCAGCAC CGGGCGTGGC TAGCGGGAAG
-----
1501  CCAACAGTTA CGCAGCCTGA ATGGCGAATG GCGCTTTGCC TGGTTTCCGG
     GGTGTGCAAT GCGTCGGACT TACCGCTTAC CGCGAAACGG ACCAAAGGCG
-----
1551  CACCAGAAGC GGTGCCGGAA AGCTGGCTGG AGTGCGATCT TCCTGAGGCC
     GTGGTCTTCG CCACGGCCTT TCGACCGACC TCACGCTAGA AGGACTCCGG
-----
1601  GATACTGTCG TCGTCCCCTC AAACCTGGCAG ATGCACGGTT ACGATGCGCC
     CTATGACAGC AGCAGGGGAG TTTGACCGTC TACGTGCCAA TGCTACGCGG
-----
1651  CATCTACACC AACGTGACCT ATCCCATTAC GGTCAATCCG CCGTTTGTTT
     GTAGATGTGG TTGCACTGGA TAGGGTAATG CCAGTTAGGC GGCAACAAG
-----
1701  CCACGGAGAA TCCGACGGGT TGTTACTCGC TCACATTTAA TGTGATGAA
     GGTGCCTCTT AGGCTGCCCA ACAATGAGCG AGTGTAATT ACAACTACTT
-----
1751  AGCTGGCTAC AGGAAGGCCA GACGCGAATT ATTTTGTATG GCGTTAACTC
     TCGACCGATG TCCTTCCGGT CTGCGCTTAA TAAAACTAC CGCAATTGAG
-----
1801  GGCGTTTCAT CTGTGGTGCA ACGGGCGCTG GGTCGGTTAC GGCCAGGACA
     CCGCAAAGTA GACACCACGT TGCCCGCGAC CCAGCCAATG CCGGTCCTGT
-----
1851  GTCGTTTGCC GTCTGAATTT GACCTGAGCG CATTTTTACG CGCCGGAGAA
     CAGCAAACGG CAGACTTAAA CTGGACTCGC GTAAAAATGC GCGGCCTCTT
-----
```



```
1901  AACCGCCTCG CGGTGATGGT GCTGCGCTGG AGTGACGGCA GTTATCTGGA
      TTGGCGGAGC GCCACTACCA CGACGCGACC TCACTGCCGT CAATAGACCT
-----
1951  AGATCAGGAT ATGTGGCGGA TGAGCGGCAT TTTCCGTGAC GTCTCGTTGC
      TCTAGTCCTA TACACCGCCT ACTCGCCGTA AAAGGCACTG CAGAGCAACG
-----
2001  TGCATAAACC GACTACACAA ATCAGCGATT TCCATGTTGC CACTCGCTTT
      ACGTATTTGG CTGATGTGTT TAGTCGCTAA AGGTACAACG GTGAGCGAAA
-----
2051  AATGATGATT TCAGCCGCGC TGTACTGGAG GCTGAAGTTC AGATGTGCGG
      TTACTACTAA AGTCGGCGCG ACATGACCTC CGACTTCAAG TCTACACGCC
-----
2101  CGAGTTGCGT GACTACCTAC GGGTAACAGT TTCTTTATGG CAGGGTGAAA
      GCTCAACGCA CTGATGGATG CCCATTGTCA AAGAAATACC GTCCCACTTT
-----
2151  CGCAGGTGCG CAGCGGCACC GCGCCTTTTC GCGGTGAAAT TATCGATGAG
      GCGTCCAGCG GTCGCCGTGG CGCGGAAAGC CGCCACTTTA ATAGCTACTC
-----
2201  CGTGGTGGTT ATGCCGATCG CGTCACACTA CGTCTGAACG TCGAAAACCC
      GCACCACCAA TACGGCTAGC GCAGTGTGAT GCAGACTTGC AGCTTTTGGG
-----
2251  GAAACTGTGG AGCGCCGAAA TCCCGAATCT CTATCGTGCG GTGGTTGAAC
      CTTTGACACC TCGCGGCTTT AGGGCTTAGA GATAGCACGC CACCAACTTG
-----
2301  TGCACACCGC CGACGGCACG CTGATTGAAG CAGAAGCCTG CGATGTCGGT
      ACGTGTGGCG GCTGCCGTGC GACTAACTTC GTCTTCGGAC GCTACAGCCA
-----
2351  TTCCGCGAGG TGCGGATTGA AAATGGTCTG CTGCTGCTGA ACGGCAAGCC
      AAGGCGCTCC ACGCCTAACT TTTACCAGAC GACGACGACT TGCCGTTTCG
-----
2401  GTTGCTGATT CGAGGCGTTA ACCGTCACGA GCATCATCCT CTGCATGGTC
      CAACGACTAA GCTCCGCAAT TGGCAGTGCT CGTAGTAGGA GACGTACCAG
-----
2451  AGGTCATGGA TGAGCAGACG ATGGTGCAGG ATATCCTGCT GATGAAGCAG
      TCCAGTACCT ACTCGTCTGC TACCACGTCC TATAGGACGA CTACTTCGTC
-----
2501  AACAACTTTA ACGCCGTGCG CTGTTTCGCAT TATCCGAACC ATCCGCTGTG
      TTGTTGAAAT TGCGGCACGC GACAAGCGTA ATAGGCTTGG TAGGCGACAC
-----
2551  GTACACGCTG TGCGACCGCT ACGGCCTGTA TGTGGTGGAT GAAGCCAATA
      CATGTGCGAC ACGCTGGCGA TGCCGGACAT ACACCACCTA CTTCCGGTTAT
-----
2601  TTGAAACCCA CGGCATGGTG CCAATGAATC GTCTGACCGA TGATCCGCGC
      AACTTTGGGT GCCGTACCAC GGTTACTTAG CAGACTGGCT ACTAGGCGCG
-----
2651  TGGCTACCGG CGATGAGCGA ACGCGTAACG CGAATGGTGC AGCGCGATCG
      ACCGATGGCC GCTACTCGCT TGCGCATTGC GCTTACCACG TCGCGCTAGC
-----
2701  TAATCACCCG AGTGTGATCA TCTGGTCGCT GGGGAATGAA TCAGGCCACG
      ATTAGTGGGC TCACACTAGT AGACCAGCGA CCCCTTACTT AGTCCGGTGC
-----
2751  GCGCTAATCA CGACGCGCTG TATCGCTGGA TCAAATCTGT CGATCCTTCC
      CGCGATTAGT GCTGCGCGAC ATAGCGACCT AGTTTAGACA GCTAGGAAGG
-----
2801  CGCCCGGTGC AGTATGAAGG CGGCGGAGCC GACACCACGG CCACCGATAT
      GCGGGCCACG TCATACTTCC GCCGCCTCGG CTGTGGTGCC GGTGGCTATA
-----
```

2851 TATTTGCCCCG ATGTACGCGC GCGTGGATGA AGACCAGCCC TTCCCGGCTG  
ATAAACGGGC TACATGCGCG CGCACCTACT TCTGGTCGGG AAGGGCCGAC  
-----  
2901 TGCCGAAATG GTCCATCAAA AAATGGCTTT CGCTACCTGG AGAGACGCGC  
ACGGCTTTTAC CAGGTAGTTT TTTACCGAAA GCGATGGACC TCTCTGCGCG  
-----  
2951 CCGCTGATCC TTTGCGAATA CGCCCACGCG ATGGGTAACA GTCTTGCGCG  
GGCGACTAGG AAACGCTTAT GCGGGTGCGC TACCCATTGT CAGAACCGCC  
-----  
3001 TTTCGCTAAA TACTGGCAGG CGTTTCGTCA GTATCCCCGT TTACAGGGCG  
AAAGCGATTT ATGACCGTCC GCAAAGCAGT CATAGGGGCA AATGTCCCGC  
-----  
3051 GCTTCGTCTG GGAAGGCTG GATCAGTCGC TGATTAAATA TGATGAAAAC  
CGAAGCAGAC CCTGACCCAC CTAGTCAGCG ACTAATTTAT ACTACTTTTG  
-----  
3101 GGCAACCCGT GGTTCGGCTTA CGGCGGTGAT TTTGGCGATA CGCCGAACGA  
CCGTGGGGCA CCAGCCGAAT GCCGCCACTA AAACCGCTAT GCGGCTTGCT  
-----  
3151 TCGCCAGTTC TGTATGAACG GTCTGGTCTT TGCCGACCGC ACGCCGCATC  
AGCGGTCAAG ACATACTTGC CAGACCAGAA ACGGCTGGCG TGCGGCGTAG  
-----  
3201 CAGCGCTGAC GGAAGCAAAA CACCAGCAGC AGTTTTTCCA GTTCCGTTTA  
GTGCGGACTG CCTTCGTTTT GTGGTCGTCG TCAAAAAGGT CAAGGCAAAAT  
-----  
3251 TCCGGGCAAA CCATCGAAGT GACCAGCGAA TACCTGTTCC GTCATAGCGA  
AGGCCCCGTTT GGTAGCTTCA CTGGTCGCTT ATGGACAAGG CAGTATCGCT  
-----  
3301 TAACGAGCTC CTGCACTGGA TGGTGGCGCT GGATGGTAAG CCGCTGGCAA  
ATTGCTCGAG GACGTGACCT ACCACCGCGA CCTACCATTG GGCACCGTT  
-----  
3351 GCGGTGAAGT GCCTCTGGAT GTCGCTCCAC AAGGTAAACA GTTGATTGAA  
CGCCACTTCA CGGAGACCTA CAGCGAGGTG TTCCATTGT CAACTAACTT  
-----  
3401 CTGCCTGAAC TACCGCAGCC GGAGAGCGCC GGGCAACTCT GGCTCACAGT  
GACGGACTTG ATGGCGTCGG CCTCTCGCGG CCCGTTGAGA CCGAGTGTC  
-----  
3451 ACGCGTAGTG CAACCGAACG CGACCGCATG GTCAGAAGCC GGGCACATCA  
TGCGCATCAC GTTGGCTTGC GCTGGCGTAC CAGTCTTCGG CCCGTGTAGT  
-----  
3501 GCGCCTGGCA GCAGTGGCGT CTGGCGGAAA ACCTCAGTGT GACGCTCCCC  
CGCGGACCGT CGTCACCGCA GACCGCCTTT TGGAGTCACA CTGCGAGGGG  
-----  
3551 GCCGCGTCCC ACGCCATCCC GCATCTGACC ACCAGCGAAA TGGATTTTTG  
CGGCGCAGGG TGCGGTAGGG CGTAGACTGG TGGTCGCTTT ACCTAAAAAC  
-----  
3601 CATCGAGCTG GGTAATAAGC GTTGGAATT TAACCGCCAG TCAGGCTTTC  
GTAGCTCGAC CCATTATTG CAACCGTTAA ATTGGCGGTC AGTCCGAAAG  
-----  
3651 TTTCACAGAT GTGGATTGGC GATAAAAAAC AACTGCTGAC GCCGCTGCGC  
AAAGTGCTA CACCTAACCG CTATTTTTTG TTGACGACTG CGGCGACGCG  
-----  
3701 GATCAGTTCA CCCGTGCACC GCTGGATAAC GACATTGGCG TAAGTGAAGC  
CTAGTCAAGT GGGCACGTGG CGACCTATTG CTGTAACCGC ATTCACTTCG  
-----  
3751 GACCCGCATT GACCCTAACG CCTGGGTCGA ACGCTGGAAG GCGGCGGGCC  
CTGGGCGTAA CTGGGATTGC GGACCCAGCT TGCGACCTTC CGCCGCCCCG

```
3801 ATTACCAGGC CGAAGCAGCG TTGTTGCAGT GCACGGCAGA TACACTTGCT
    TAATGGTCCG GCTTCGTCGC AACAACTCA CGTGCCGTCT ATGTGAACGA
-----
3851 GATGCGGTGC TGATTACGAC CGCTCACGCG TGGCAGCATC AGGGGAAAAC
    CTACGCCACG ACTAATGCTG GCGAGTGCGC ACCGTCGTAG TCCCCTTTTG
-----
3901 CTTATTTATC AGCCGGAAAA CCTACCGGAT TGATGGTAGT GGTCAAATGG
    GAATAAATAG TCGGCCTTTT GGATGGCCTA ACTACCATCA CCAGTTTACC
-----
3951 CGATTACCGT TGATGTTGAA GTGGCGAGCG ATACACCGCA TCCGGCGCGG
    GCTAATGGCA ACTACAACIT CACCGCTCGC TATGTGGCGT AGGCCGCGCC
-----
4001 ATTGGCCTGA ACTGCCAGCT GGCGCAGGTA GCAGAGCGGG TAAACTGGCT
    TAACCGGACT TGACGGTCGA CCGCGTCCAT CGTCTCGCCC ATTTGACCGA
-----
4051 CGGATTAGGG CCGCAAGAAA ACTATCCCGA CCGCCTTACT GCCGCCTGTT
    GCCTAATCCC GCGTTTCTTT TGATAGGGCT GCGGGAATGA CGGCGGACAA
-----
4101 TTGACCGCTG GGATCTGCCA TTGTCAGACA TGTATACCCC GTACGTCTTC
    AACTGGCGAC CCTAGACGGT AACAGTCTGT ACATATGGGG CATGCAGAAG
-----
4151 CCGAGCGAAA ACGGTCTGCG CTGCGGGACG CGCGAATTGA ATTATGGCCC
    GGCTCGCTTT TGCCAGACGC GACGCCCTGC GCGCTTAACT TAATACCGGG
-----
4201 ACACCAGTGG CGCGGCGACT TCCAGTTCAA CATCAGCCGC TACAGTCAAC
    TGTGGTCACC GCGCCGCTGA AGGTCAAGTT GTAGTCGGCG ATGTCAGTTG
-----
4251 AGCAACTGAT GGAAACCAGC CATCGCCATC TGCTGCACGC GGAAGAAGGC
    TCGTTGACTA CCTTTGGTCG GTAGCGGTAG ACGACGTGCG CCTTCTTCCG
-----
4301 ACATGGCTGA ATATCGACGG TTTCCATATG GGGATTGGTG GCGACGACTC
    TGTACCGACT TATAGCTGCC AAAGGTATAC CCCTAACCAC CGCTGCTGAG
-----
4351 CTGGAGCCCC TCAGTATCGG CGGAATTCCA GCTGAGCGCC GGTCGCTACC
    GACCTCGGGC AGTCATAGCC GCCTTAAGGT CGACTCGCGG CCAGCGATGG
-----
4401 ATTACCAATT GGTCTGGTGT CAAAAAAGAT CTGGAGGTGG TGGCAGCAGG
    TAATGGTCAA CCAGACCACA GTTTTTTCTA GACCTCCACC ACCGTCGTCC
-----
4451 CCTTGGCGCG CCGGATCCTT AATTAACAAT TGACCGGTAA TAATAGGTAG
    GGAACCGCGC GGCCTAGGAA TTAATTGTTA ACTGGCCATT ATTATCCATC
-----
4501 ATAAGTGAAT GATTAGATGC ATTGATCCCT CGACCAATTC CGGTTATTTT
    TATTCACCTG CTAATCTACG TAACTAGGGA GCTGGTTAAG GCCAATAAAA
-----
4551 CCACCATATT GCCGTCTTTT GGCAATGTGA GGGCCCGGAA ACCTGGCCCT
    GGTGGTATAA CCGCAGAAAA CCGTTACACT CCCGGGCCTT TGGACCGGGA
-----
4601 GTCTTCTTGA CGAGCATTCC TAGGGGTCTT TCCCCTCTCG CCAAAGGAAT
    CAGAAGAACT GCTCGTAAGG ATCCCAGAA AGGGGAGAGC GGTTTCCTTA
-----
4651 GCAAGGTCTG TTGAATGTCG TGAAGGAAGC AGTTCCTCTG GAAGCTTCTT
    CGTTCAGAC AACTTACAGC ACTTCCTTCG TCAAGGAGAC CTTCAAGAA
-----
4701 GAAGACAAAC AACGTCTGTA GCGACCCTTT GCAGGCAGCG GAACCCCCCA
    CTTCTGTTTG TTGCAGACAT CGCTGGGAAA CGTCCGTCGC CTTGGGGGGT
-----
```

```
4751 CCTGGCGACA GGTGCCTCTG CGGCCAAAAG CCACGTGTAT AAGATACACC
    GGACCGCTGT CCACGGAGAC GCCGGTTTTT GGTGCACATA TTCTATGTGG
-----
4801 TGCAAAGGCG GCACAACCCC AGTGCCACGT TGTGAGTTGG ATAGTTGTGG
    ACGTTTCCGC CGTGTGTTGGG TCACGGTGCA ACACTCAACC TATCAACACC
-----
4851 AAAGAGTCAA ATGGCTCTCC TCAAGCGTAT TCAACAAGGG GCTGAAGGAT
    TTTCTCAGTT TACCGAGAGG AGTTTCGCATA AGTTGTTCCC CGACTTCCTA
-----
4901 GCCCAGAAGG TACCCCATTT TATGGGATCT GATCTGGGGC CTCGGTGCAC
    CGGGTCTTCC ATGGGGTAAC ATACCCTAGA CTAGACCCCG GAGCCACGTG
-----
4951 ATGCTTTTACA TGTGTTTAGT CGAGGTAAAA AAACGTCTAG GCCCCCGAA
    TACGAAATGT ACACAAATCA GCTCCAATTT TTTGCAGATC CGGGGGGCTT
-----
5001 CCACGGGGAC GTGGTTTTTC TTTGAAAAAC ACGATGATAA TACCATGATT
    GGTGCCCTTG CACCAAAAGG AAACTTTTTG TGCTACTATT ATGGTACTAA
-----
5051 GAACAAGATG GATTGCACGC AGGTTCTCCG GCCGCTTGGG TGGAGAGGCT
    CTTGTTCTAC CTAACGTGCG TCCAAGAGGC CGGCGAACCC ACCTCTCCGA
-----
5101 ATTCGGCTAT GACTGGGCAC AACAGACAAT CGGCTGCTCT GATGCCGCCG
    TAAGCCGATA CTGACCCGTG TTGTCTGTTA GCCGACGAGA CTACGGCGGC
-----
5151 TGTTCGGGCT GTCAGCGCAG GGGCGCCCGG TTCTTTTTGT CAAGACCGAC
    ACAAGGCCGA CAGTCGCGTC CCCGCGGGCC AAGAAAAACA GTTCTGGCTG
-----
5201 CTGTCCGGTG CCCTGAATGA ACTGCAGGAC GAGGCAGCGC GGCTATCGTG
    GACAGGCCAC GGGACTTACT TGACGTCCTG CTCCGTCGCG CCGATAGCAC
-----
5251 GCTGGCCACG ACGGGCGTTC CTTGCGCAGC TGTGCTCGAC GTTGTCACTG
    CGACCGGTGC TGCCCGCAAG GAACGCGTCG ACACGAGCTG CAACAGTGAC
-----
5301 AAGCGGGAAG GGACTGGCTG CTATTGGGCG AAGTGCCGGG GCAGGATCTC
    TTCGCCCTTC CCTGACCGAC GATAACCCGC TTCACGGCCC CGTCCTAGAG
-----
5351 CTGTCATCTC ACCTTGCTCC TGCCGAGAAA GTATCCATCA TGGCTGATGC
    GACAGTAGAG TGGAACGAGG ACGGCTCTTT CATAGGTAGT ACCGACTACG
-----
5401 AATGCGGCGG CTGCATACGC TTGATCCGGC TACCTGCCCA TTCGACCACC
    TTACGCCGCC GACGTATGCG AACTAGGCCG ATGGACGGGT AAGCTGGTGG
-----
5451 AAGCGAAACA TCGCATCGAG CGAGCACGTA CTCGGATGGA AGCCGGTCTT
    TTCGCTTTGT AGCGTAGCTC GCTCGTGCAT GAGCCTACCT TCGGCCAGAA
-----
5501 GTCGATCAGG ATGATCTGGA CGAAGAGCAT CAGGGGCTCG CGCCAGCCGA
    CAGCTAGTCC TACTAGACCT GCTTCTCGTA GTCCCGGAGC GCGGTCGGCT
-----
5551 ACTGTTCCGCC AGGCTCAAGG CGCGCATGCC CGACGGCGAG GATCTCGTCC
    TGACAAGCGG TCCGAGTTCC GCGCGTACGG GCTGCCGCTC CTAGAGCAGC
-----
5601 TGACCCATGG CGATGCCTGC TTGCCGAATA TCATGGTGGA AAATGGCCGC
    ACTGGGTACC GCTACGGACG AACGGCTTAT AGTACCACCT TTTACCGGCG
-----
5651 TTTTCTGGAT TCATCGACTG TGGCCGGCTG GGTGTGGCGG ACCGCTATCA
    AAAAGACCTA AGTAGCTGAC ACCGGCCGAC CCACACCGCC TGGCGATAGT
-----
```

```
5701  GGACATAGCG TTGGCTACCC GTGATATTGC TGAAGAGCTT GGCGGCGAAT
      CCTGTATCGC AACCGATGGG CACTATAACG ACTTCTCGAA CCGCCGCTTA
-----
5751  GGGCTGACCG CTTCTCGTG CTTTACGGTA TCGCCGCTCC CGATTTCGAG
      CCCGACTGGC GAAGGAGCAC GAAATGCCAT AGCGGCGAGG GCTAAGCGTC
-----
5801  CGCATCGCCT TCTATCGCCT TCTTGACGAG TTCTTCTGAG CGGGACTCTG
      GCGTAGCGGA AGATAGCGGA AGAACTGCTC AAGAAGACTC GCCCTGAGAC
-----
5851  GGGTTTCGCAT CGATAAAATA AAAGATTTTA TTTAGTCTCC AGAAAAAGGG
      CCCAAGCGTA GCTATTTTAT TTTCTAAAT AAATCAGAGG TCTTTTCC
-----
5901  GGAATGAAA GACCCACCT GTAGGTTTGG CAAGCTAGCT TAAGTAACGC
      CCCTTACTTT CTGGGGTGA CATCCAAACC GTTCGATCGA ATTCATTGCG
-----
5951  CATTTTGCAA GGCATGGAAA AATACATAAC TGAGAATAGA GAAGTTCAGA
      GTAAACGTT CCGTACCTTT TTATGTATTG ACTCTTATCT CTTCAAGTCT
-----
6001  TCAAGGTCAG GAACAGATGG AACAGCTGAA TATGGGCCAA ACAGGATATC
      AGTTCAGTC CTTGTCTACC TTGTCGACTT ATACCCGGTT TGTCTATAG
-----
6051  TGTGGTAAGC AGTTCCTGCC CCGGCTCAGG GCCAAGAACA GATGGAACAG
      ACACCATTTC TCAAGGACGG GGCCGAGTCC CGGTTCTTGT CTACCTTGTC
-----
6101  CTGAATATGG GCCAAACAGG ATATCTGTGG TAAGCAGTTC CTGCCCCGGC
      GACTTATACC CGGTTTGTCC TATAGACACC ATTCGTCAAG GACGGGGCCG
-----
6151  TCAGGGCCAA GAACAGATGG TCCCAGATG CGGTCCAGCC CTCAGCAGTT
      AGTCCCGGTT CTTGTCTACC AGGGGTCTAC GCCAGGTCGG GAGTCGTCAA
-----
6201  TCTAGAGAAC CATCAGATGT TTCCAGGGTG CCCAAGGAC CTGAAATGAC
      AGATCTCTTG GTAGTCTACA AAGGTCCAC GGGGTTCTTG GACTTTACTG
-----
6251  CCTGTGCCTT ATTTGAACTA ACCAATCAGT TCGCTTCTCG CTTCTGTTTG
      GGACACGGAA TAACTTGAT TGGTTAGTCA AGCGAAGAGC GAAGACAAGC
-----
6301  CGCGCTTCTG CTCCCCGAGC TCAATAAAAG AGCCACAAC CCCTCACTCG
      GCGCGAAGAC GAGGGGCTCG AGTTATTTTC TCGGGTGTG GGGAGTGAGC
-----
6351  GGGCGCCAGT CCTCCGATTG ACTGAGTCGC CCGGGTACCC GTGTATCCAA
      CCCGCGGTCA GGAGGCTAAC TGAATCAGCG GGCCATGGG CACATAGGTT
-----
6401  TAAACCCTCT TGCAGTTGCA TCCGACTTGT GGTCTCGCTG TTCCTTGGA
      ATTTGGGAGA ACGTCAACGT AGGCTGAACA CCAGAGCGAC AAGGAACCCT
-----
6451  GGGTCTCCTC TGAGTGATTG ACTACCCGTC AGCGGGGGTC TTTCATTCAT
      CCCAGAGGAG ACTCACTAAC TGATGGGCAG TCGCCCCCAG AAAGTAAGTA
-----
6501  GCAGCATGTA TCAAAATTAA TTTGGTTTTT TTTCTTAAGT ATTTACATTA
      CGTCGTACAT AGTTTTAATT AAACCAAAAA AAAGAATTCA TAAATGTAAT
-----
6551  AATGGCCATA GTTGCAATTAA TGAATCGGCC AACGCGCGGG GAGAGGCGGT
      TTACCGGTAT CAACGTAATT ACTTAGCCGG TTGCGCGCCC CTCTCCGCCA
-----
6601  TTGCGTATTG GCGCTCTTCC GCTTCCTCGC TCACTGACTC GCTGCGCTCG
      AACGCATAAC CGCGAGAAGG CGAAGGAGCG AGTGAATGAG CGACGCGAGC
-----
```

```
6651  GTCGTTTCGGC TCGGGCGAGC GGTATCAGCT CACTCAAAGG CGGTAATACG
      CAGCAAGCCG ACGCCGCTCG CCATAGTCGA GTGAGTTTCC GCCATTATGC
-----
6701  GTTATCCACA GAATCAGGGG ATAACGCAGG AAAGAACATG TGAGCAAAAG
      CAATAGGTGT CTTAGTCCCC TATTGCGTCC TTTCTTGAC ACTCGTTTTC
-----
6751  GCCAGCAAAA GGCCAGGAAC CGTAAAAAGG CCGCGTTGCT GGCCTTTTTT
      CCGTCGTTTT CCGGTCCTTG GCATTTTTTC GGCACAACGA CCGCAAAAAG
-----
6801  CATAGGCTCC GCCCCCTGA CGAGCATCAC AAAAATCGAC GCTCAAGTCA
      GTATCCGAGG CGGGGGGACT GCTCGTAGTG TTTTITAGCTG CGAGTTCAGT
-----
6851  GAGGTGGCGA AACCCGACAG GACTATAAAG ATACCAGGCG TTTCCCCCTG
      CTCCACCCTG TTGGGCTGTC CTGATATTTT TATGGTCCGC AAAGGGGGAG
-----
6901  GAAGCTCCCT CGTGCGCTCT CCTGTTCCGA CCCTGCCGCT TACCGGATAC
      CTTTCGAGGA GCACGCGAGA GGACAAGGCT GGGACGGCGA ATGGCCTATG
-----
6951  CTGTCCGCCT TTCTCCCTTC GGGAAGCGTG GCGCTTTTCT ATAGCTCACG
      GACAGGCGGA AAGAGGGAAG CCCTTCGCAC CGCGAAAGAG TATCGAGTGC
-----
7001  CTGTAGGTAT CTCAGTTCGG TGTAGGTCGT TCGCTCCAAG CTGGGCTGTG
      GACATCCATA GAGTCAAGCC ACATCCAGCA AGCGAGGTTT GACCCGACAC
-----
7051  TGCACGAACC CCCCCTTCAG CCCGACCGCT GCGCCTTATC CGGTAAGTAT
      ACGTGCTTGG GGGGCAAGTC GGGCTGGCGA CGCGGAATAG GCCATTGATA
-----
7101  CGTCTTGAGT CCAACCCGGT AAGACACGAC TTATCGCCAC TGGCAGCAGC
      GCAGAACTCA GGTTGGGCCA TTCTGTGCTG AATAGCGGTG ACCGTCGTCG
-----
7151  CACTGGTAAC AGGATTAGCA GAGCGAGGTA TGTAGGCGGT GCTACAGAGT
      GTGACCATTG TCCTAATCGT CTCGCTCCAT ACATCCGCCA CGATGTCTCA
-----
7201  TCTTGAAGTG GTGGCCTAAC TACGGCTACA CTAGAAGAAC AGTATTTGGT
      AGAACTTCAC CACCGGATTG ATGCCGATGT GATCTTCTTG TCATAAACCA
-----
7251  ATCTGCGCTC TGCTGAAGCC AGTTACCTTC GGAAAAAGAG TTGGTAGCTC
      TAGACGCGAG ACGACTTCGG TCAATGGAAG CCTTTTCTC AACCATCGAG
-----
7301  TTGATCCGGC AAACAAACCA CCGCTGGTAG CGGTGGTTTT TTTGTTTGCA
      AACTAGGCCG TTTGTTTGGT GGCGACCATC GCCACCAAAA AAACAAACGT
-----
7351  AGCAGCAGAT TACGCGCAGA AAAAAAGGAT CTCAAGAAGA TCCTTTGATC
      TCGTCGTCTA ATGCGCGTCT TTTTTCCTA GAGTCTTCT AGGAAACTAG
-----
7401  TTTTCTACGG GGTCTGACGC TCAGTGAAC GAAAACTCAC GTTAAGGGAT
      AAAAGATGCC CCAGACTGCG AGTCACCTTG CTTTGTAGTG CAATTCCCTA
-----
7451  TTTGGTCATG AGATTATCAA AAAGGATCTT CACCTAGATC CTTTTCGGC
      AAACCAGTAC TCTAATAGTT TTTCTAGAA GTGGATCTAG GAAAACGCCG
-----
7501  CGCAAATCAA TCTAAAGTAT ATATGAGTAA ACTTGGTCTG ACAGTTACCA
      GCGTTTAGTT AGATTTTATA TATACTCATT TGAACCAGAC TGTCAATGGT
-----
7551  ATGCTTAATC AGTGAGGCAC CTATCTCAGC GATCTGTCTA TTTCGTTTAT
      TACGAATTAG TCACTCCGTG GATAGAGTCG CTAGACAGAT AAAGCAAGTA
-----
```

7601 CCATAGTTGC CTGACTCCCC GTCGTGTAGA TAACTACGAT ACGGGAGGGC  
GGTATCAACG GACTGAGGGG CAGCACATCT ATTGATGCTA TGCCCTCCCG

7651 TTACCATCTG GCCCCAGTGC TGCAATGATA CCGCGAGACC CACGCTCACC  
AATGGTAGAC CGGGGTCACG ACGTTACTAT GCGGCTCTGG GTGCGAGTGG

7701 GGCTCCAGAT TTATCAGCAA TAAACCAGCC AGCCGGAAGG GCCGAGCGCA  
CCGAGGTCTA AATAGTCGTT ATTTGGTCGG TCGGCCTTCC CGGCTCGCGT

7751 GAAGTGGTCC TGCAACTTTA TCCGCCTCCA TCCAGTCTAT TAATTGTTGC  
CTTACCAGG ACGTTGAAAT AGGCGGAGGT AGGTCAGATA ATTAACAACG

7801 CGGGAAGCTA GAGTAAGTAG TTCGCCAGTT AATAGTTTGC GCAACGTTGT  
GCCCTTCGAT CTCATTCAATC AAGCGGTCAA TTATCAAACG CGTTGCAACA

7851 TGCCATTGCT ACAGGCATCG TGGTGTACG CTCGTCGTTT GGTATGGCTT  
ACGGTAACGA TGTCCTAGC ACCACAGTGC GAGCAGCAAA CCATACCGAA

7901 CATTAGCTC CGGTTCCCAA CGATCAAGGC GAGTTACATG ATCCCCCATG  
GTAAGTCGAG GCCAAGGGTT GCTAGTTCCG CTCAATGTAC TAGGGGGTAC

7951 TTGTGCAAAA AAGCGGTTAG CTCCTTCGGT CCTCCGATCG TTGTCAGAAG  
AACACGTTTT TTCGCCAATC GAGGAAGCCA GGAGGCTAGC AACAGTCTTC

8001 TAAGTTGGCC GCAGTGTAT CACTCATGGT TATGGCAGCA CTGCATAATT  
ATTCAACCGG CGTCACAATA GTGAGTACCA ATACCGTCGT GACGTATTAA

8051 CTCTTACTGT CATGCCATCC GTAAGATGCT TTTCTGTGAC TGGTGAGTAC  
GAGAAATGACA GTACGGTAGG CATTCTACGA AAAGACACTG ACCACTCATG

8101 TCAACCAAGT CATTCTGAGA ATAGTGTATG CGGCGACCGA GTTGCTCTTG  
AGTTGGTTCA GTAAGACTCT TATCACATAC GCCGCTGGCT CAACGAGAAC

8151 CCCGGCGTCA ATACGGGATA ATACCGCGCC ACATAGCAGA ACTTTAAAAG  
GGGCCGCGT TATGCCCTAT TATGGCGCGG TGTATCGTCT TGAAATTTTC

8201 TGCTCATCAT TGGAAAACGT TCTTCGGGGC GAAAACTCTC AAGGATCTTA  
ACGAGTAGTA ACCTTTTGCA AGAAGCCCCG CTTTGTGAGAG TTCCTAGAAT

8251 CCGCTGTTGA GATCCAGTTC GATGTAACCC ACTCGTGCAC CCAACTGATC  
GGCGACAAC CTAGGTCAAG CTACATTGGG TGAGCACGTG GGTTGACTAG

8301 TTCAGCATCT TTTACTTTCA CCAGCGTTTC TGGGTGAGCA AAAACAGGAA  
AAGTCGTAGA AAATGAAAGT GGTCGCAAAG ACCCACTCGT TTTTGTCTT

8351 GGCAAAATGC CGCAAAAAAG GGAATAAGGG CGACACGGAA ATGTTGAATA  
CCGTTTTACG GCGTTTTTTC CCTTATTCCC GCTGTGCCTT TACAATTAT

8401 CTCATACTCT TCCTTTTTC ATATTATTGA AGCATTATC AGGGTTATTG  
GAGTATGAGA AGGAAAAAGT TATAATAACT TCGTAAATAG TCCCAATAAC

8451 TCTCATGAGC GGATACATAT TTGAATGTAT TTAGAAAAAT AAACAAATAG  
AGAGTACTCG CCTATGTATA AACTTACATA AATCTTTTTA TTTGTTTATC

8501 GGGTTCCGCG CACATTTT  
CCCAAGGCGC GTGTAAAG

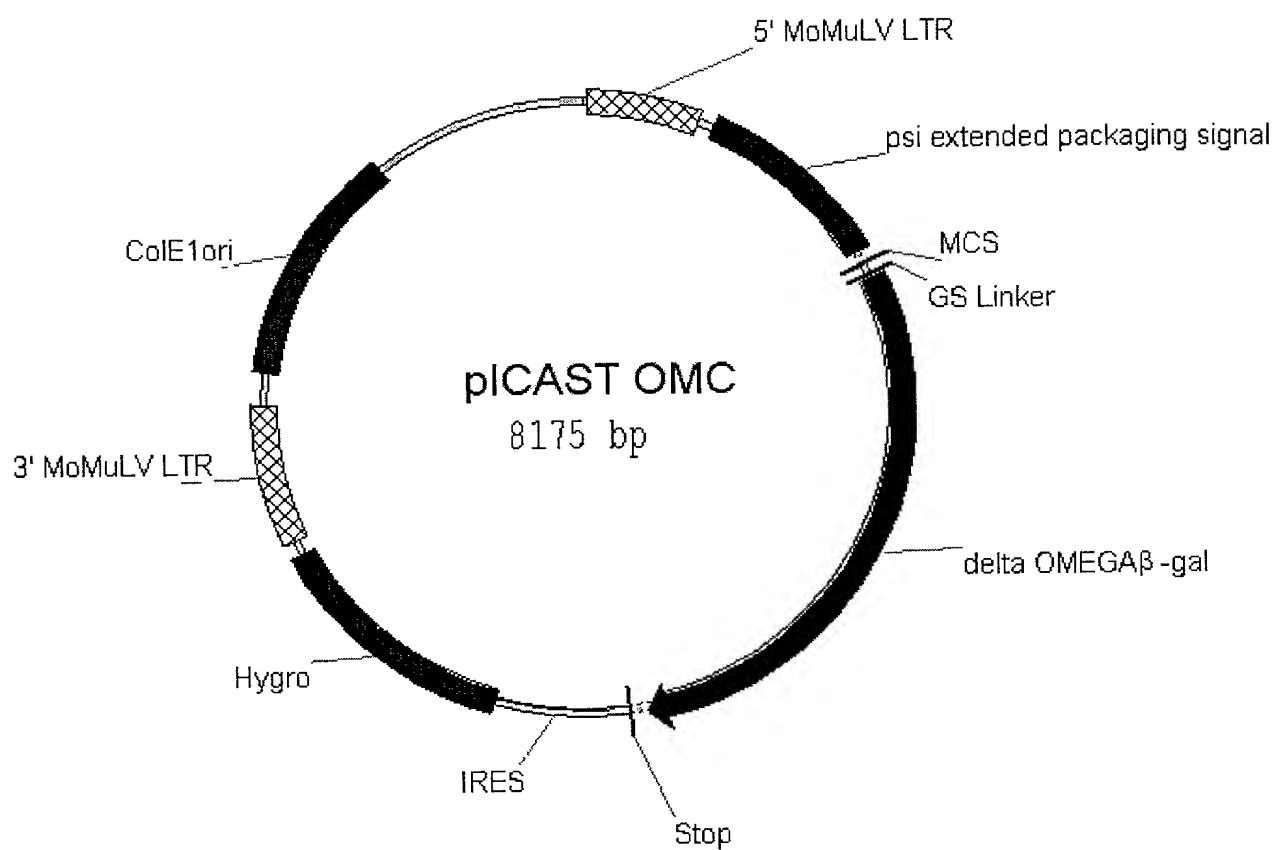


Figure 12A



```

1  CTGCAGCCTG AATATGGGCC AAACAGGATA TCTGTGGTAA GCAGTTCCTG
   GACGTCCGAC TTATACCCGG TTTGTCCTAT AGACACCATT CGTCAAGGAC
-----
51  CCCC GGCTCA GGGCCAAGAA CAGATGGAAC AGCTGAATAT GGGCCAAACA
   GGGGCCGAGT CCCGGTTCTT GTCTACCTTG TCGACTTATA CCCGGTTTGT
-----
101 GGATATCTGT GGTAAGCAGT TCCTGCCCCG GCTCAGGGCC AAGAACAGAT
   CCTATAGACA CCATTCGTCA AGGACGGGGC CGAGTCCCGG TTCTTGCTA
-----
151 GGTCCCCAGA TGC GGTTCCAG CCTCAGCAG TTTCTAGAGA ACCATCAGAT
   CCAGGGGTCT ACGCCAGGTC GGGAGTCGTC AAAGATCTCT TGGTAGTCTA
-----
201 GTTTC CAGGG TGCCCCAAGG ACCTGAAATG ACCCTGTGCC TTATTTGAAC
   CAAAGGTCCC ACGGGGTTC TGGACTTTAC TGGGACACGG AATAAACTTG
-----
251 TAACCAATCA GTTCGCTTCT CGCTTCTGTT CGCGCGCTTC TGCTCCCCGA
   ATTGGTTAGT CAAGCGAAGA GCGAAGACAA GCGCGCGAAG ACGAGGGGCT
-----
301 GCTCAATAAA AGAGCCCACA ACCCCTCACT CGGGGCGCCA GTCCTCCGAT
   CGAGTTATTT TCTCGGGTGT TGGGGAGTGA GCCCCGCGGT CAGGAGGCTA
-----
351 TGA CTGAGTC GCCCGGGTAC CCGTGTATCC AATAAACCCCT CTTGCAGTTG
   ACTGACTCAG CGGGCCCATG GGCACATAGG TTATTTGGGA GAACGTCAAC
-----
401 CATCCGACTT GTGGTCTCGC TGTTCTTGG GAGGGTCTCC TCTGAGTGAT
   GTAGGCTGAA CACCAGAGCG ACAAGGAACC CTCCCAGAGG AGACTACTA
-----
451 TGACTACCCG TCAGCGGGGG TCTTTCATTT GGGGGCTCGT CCGGGATCGG
   ACTGATGGGC AGTCGCCCCC AGAAAGTAAA CCCCCGAGCA GGCCCTAGCC
-----
501 GAGACCCCTG CCCAGGGACC ACCGACCCAC CACCGGGAGG CAAGCTGGCC
   CTCTGGGGAC GGGTCCCTGG TGGCTGGGTG GTGGCCCTCC GTTCGACCGG
-----
551 AGCAACTTAT CTGTGTCTGT CCGATTGTCT AGTGTCTATG ACTGATTTTA
   TCGTTGAATA GACACAGACA GGCTAACAGA TCACAGATAC TGA TAAAT
-----
601 TGCGCCTGCG TCGGTACTAG TTAGCTAACT AGCTCTGTAT CTGGCGGACC
   ACGCGGACGC AGCCATGATC AATCGATTGA TCGAGACATA GACCGCCTGG
-----
651 CGTGGTGGA CTGACGAGTT CTGAACACCC GGCCGCAACC CTGGGAGACG
   GCACCACCTT GACTGCTCAA GACTTGTGGG CCGGCGTTGG GACCCTCTGC
-----
701 TCCCAGGGAC TTTGGGGGCC GTTTTGTGG CCCGACCTGA GGAAGGGAGT
   AGGGTCCCTG AAACCCCGG CAAAAACACC GGGCTGGACT CCTTCCTCA
-----
751 CGATGTGGAA TCCGACCCCG TCAGGATATG TGGTTCTGGT AGGAGACGAG
   GCTACACCTT AGGCTGGGGC AGTCCTATAC ACCAAGACCA TCCTCTGCTC
-----
801 AACCTAAAAC AGTTCCCGCC TCCGTCTGAA TTTTGTCTT CGGTTTGGAA
   TTGGATTTTG TCAAGGGCGG AGGCAGACTT AAAACGAAA GCCAAACCTT
-----
851 CCGAAGCCGC GCGTCTTGTC TGCTGCAGCA TCGTTCTGTG TTGTCTCTGT
   GGCTTCGGCG CGCAGAACAG ACGACGTCGT AGCAAGACAC AACAGAGACA
-----
901 CTGACTGTGT TTCTGTATTT GTCTGAAAAT TAGGGCCAGA CTGTTACCAC
   GACTGACACA AAGACATAAA CAGACTTTTA ATCCCGGTCT GACAATGGTG
-----

```

FIGURE 12B

```

951  TCCCTTAAGT TTGACCTTAG GTAAC TGGAA AGATGTCGAG CGGCTCGCTC
     AGGGAATTCA AACTGGAATC CATTGACCTT TCTACAGCTC GCCGAGCGAG
-----
1001 ACAAC CAGTC GGTAGATGTC AAGAAGAGAC GTTGGGTTAC CTTCTGCTCT
     TGTGTGGTCAG CCATCTACAG TTCTTCTCTG CAACCCAATG GAAGACGAGA
-----
1051 GCAGAAATGGC CAACCTTTTAA CGTCGGATGG CCGCGAGACG GCACCTTTTAA
     CGTCTTACCG GTTGGAATTT GCAGCCTACC GGCCTCTCTG CGTGGAATTT
-----
1101 CCGAGACCTC ATCACCAGG TTAAGATCAA GGTCTTTTCA CCTGGCCCGC
     GGCTCTGGAG TAGTGGGTCC AATTCTAGTT CCAGAAAAGT GGACCGGGCG
-----
1151 ATGGACACCC AGACCAGGTC CCCTACATCG TGACCTGGGA AGCCTTGGCT
     TACCTGTGGG TCTGGTCCAG GGGATGTAGC ACTGGACCCT TCGGAACCGA
-----
1201 TTTGACCCCC CTCCCTGGGT CAAGCCCTTT GTACACCCTA AGCCTCCGCC
     AAAC TGGGGG GAGGGACCCA GTTCGGGAAA CATGTGGGAT TCGGAGGCGG
-----
1251 TCCTCTTCTT CCATCCGCCC CGTCTCTCCC CTTTGAACCT CCTCGTTCTGA
     AGGAGAAGGA GGTAGGCGGG GCAGAGAGGG GGAACCTGGA GGAGCAAGCT
-----
1301 CCGCGCTCTG ATCCTCCCTT TATCCAGCCC TCACTCCTTC TCTAGGCGCC
     GGGGCGGAGC TAGGAGGGAA ATAGGTCGGG AGTGAGGAAG AGATCCGCGG
-----
1351 GGCCGCTCTA GCCCATTAA TACGACTCACT ATAGGGCGAT TCGAATCAGG
     CCGGCGAGAT CGGGTAATTA TGCTGAGTGA TATCCCGCTA AGCTTAGTCC
-----
1401 CCTTGGCGCG CCGGATCCTT AATTAAGCGC AATTGGGAGG TGGCGGTAGC
     GGAACCGCGC GGCCTAGGAA TTAATTCGCG TTAACCTTCC ACCGCCATCG
-----
1451 CTCGAGATGG GCGTGATTAC GGATTCACTG GCCGTCGTTT TACAACGTCG
     GAGCTCTACC CGCACTAATG CCTAAGTGAC CGGCAGCAA ATGTTGCAGC
-----
1501 TGA TGGGAA AACCTGGCG TTACCCAACT TAATCGCCTT GCAGCACATC
     ACTGACCCTT TTGGGACCGC AATGGGTTGA ATTAGCGGAA CGTCGTGTAG
-----
1551 CCCCTTTTCG CAGCTGGCGT AATAGCGAAG AGGCCCGCAC CGATCGCCCT
     GGGGAAAGCG GTCGACCGCA TTATCGCTTC TCCGGGCGTG GCTAGCGGGA
-----
1601 TCCCAACAGT TACG CAGCCT GAATGGCGAA TGGCGCTTTG CCTGGTTTCC
     AGGGTTGTCA ATGCGTCGGA CTTACCGCTT ACCGCGAAAC GGACCAAAGG
-----
1651 GGCACCAGAA GCGGTGCCCG AAAGCTGGCT GGAGTGCGAT CTTCTTGAGG
     CCGTGGTCTT CGCCACGGCC TTTGACCGA CCTCAGCTA GAAGGACTCC
-----
1701 CCGATACTGT CGTCGTCCCC TCAAAC TGGC AGATGCACGG TTACGATGCG
     GGCTATGACA GCAGCAGGGG AGTTTGACCG TCTACGTGCC AATGCTACGC
-----
1751 CCCATCTACA CCAACGTGAC CTATCCCAT AC GGTC AATC CGCCGTTTGT
     GGGTAGATGT GGTGCACTG GATAGGGTAA TGCCAGTTAG GCGGCAAACA
-----
1801 TCCCACGGAG AATCCGACGG GTTGTTACTC GCTCACATTT AATGTTGATG
     AGGGTGCCCT TTAGGCTGCC CAACAATGAG CGAGTGTAAT TTACAACCTAC
-----
1851 AAAGCTGGCT ACAGGAAGGC CAGACGCGAA TTATTTTGA TGGCGTTAAC
     TTTGACCGA TGCTCTCCG GTCTGCGCTT AATAAAACT ACCGCAATTG
-----

```

1901 TCGGCGTTTC ATCTGTGGTG CAACGGGCGC TGGGTCGGTT ACGGCCAGGA  
AGCCGCAAAG TAGACACCAC GTTGCCCGCG ACCCAGCCAA TGCCGGTCCT

1951 CAGTCGTTTG CCGTCTGAAT TTGACCTGAG CGCATTTTTA CGCGCCGGAG  
GTCAGCAAAC GGCAGACTTA AACTGGACTC GCGTAAAAAT GCGCGGCCTC

2001 AAAACCGCCT CGCGGTGATG GTGCTGCGCT GGAGTGACGG CAGTTATCTG  
TTTTGGCGGA GCGCCACTAC CACGACGCGA CCTCACTGCC GTCAATAGAC

2051 GAAGATCAGG ATATGTGGCG GATGAGCGGC ATTTTCCGTG ACGTCTCGTT  
CTTCTAGTCC TATACACCGC CTACTCGCCG TAAAAGGCAC TGCAGAGCAA

2101 GCTGCATAAA CCGACTACAC AAATCAGCGA TTTCCATGTT GCCACTCGCT  
CGACGTATTT GGCTGATGTG TTTAGTCGCT AAAGGTACAA CGGTGAGCGA

2151 TTAATGATGA TTTCAGCCGC GCTGTACTGG AGGCTGAAGT TCAGATGTGC  
AATTACTACT AAAGTCGGCG CGACATGACC TCCGACTTCA AGTCTACACG

2201 GCGGAGTTGC GTGACTACCT ACGGGTAACA GTTCTTTTAT GGCAGGGTGA  
CCGCTCAACG CACTGATGGA TGCCCATTGT CAAAGAAATA CCGTCCCACT

2251 AACGCAGGTC GCCAGCGGCA CCGCGCCTTT CGGCGGTGAA ATTATCGATG  
TTGCGTCCAG CGGTCGCCGT GGC GCGGAAA GCCGCCACTT TAATAGCTAC

2301 AGCGTGGTGG TTATGCCGAT CGCGTCACAC TACGTCTGAA CGTCGAAAAC  
TCGCACCACC AATACGGCTA GCGCAGTGTG ATGCAGACTT GCAGCTTTTG

2351 CCGAAACTGT GGAGCGCCGA AATCCCGAAT CTCTATCGTG CGGTGGTTGA  
GGCTTTGACA CCTCGCGGCT TTAGGGCTTA GAGATAGCAC GCCACCAACT

2401 ACTGCACACC GCCGACGGCA CGCTGATTGA AGCAGAAGCC TGCGATGTCG  
TGACGTGTGG CGGCTGCCGT GCGACTAACT TCGTCTTCGG ACGCTACAGC

2451 GTTTCCGCGA GGTGCGGATT GAAAATGGTC TGCTGCTGCT GAACGGCAAG  
CAAAGGCGCT CCACGCCTAA CTTTACCAG ACGACGACGA CTTGCCGTTT

2501 CCGTTGCTGA TTCGAGGCGT TAACCGTCAC GAGCATCATC CTCTGCATGG  
GGCAACGACT AAGCTCCGCA ATTGGCAGTG CTCGTAGTAG GAGACGTACC

2551 TCAGGTCATG GATGAGCAGA CGATGGTGCA GGATATCCTG CTGATGAAGC  
AGTCCAGTAC CTACTCGTCT GCTACCACGT CCTATAGGAC GACTACTTCG

2601 AGAACAACCT TAACGCCGTG CGCTGTTTCG ATTATCCGAA CCATCCGCTG  
TCTTGTTGAA ATTGCGGCAC GCGACAAGCG TAATAGGCTT GGTAGGCGAC

2651 TGGTACACGC TGTGCGACCG CTACGGCCTG TATGTGGTGG ATGAAGCCAA  
ACCATGTGCG ACACGCTGGC GATGCCGGAC ATACACCACC TACTTCGGTT

2701 TATTGAAACC CACGGCATGG TGCCAATGAA TCGTCTGACC GATGATCCGC  
ATAACTTTGG GTGCCGTACC ACGGTTACTT AGCAGACTGG CTACTAGGCG

2751 GCTGGCTACC GGCGATGAGC GAACGCGTAA CGCGAATGGT GCAGCGCGAT  
CGACCGATGG CCGCTACTCG CTTGCGCATT GCGCTTACCA CGTCGCGCTA

2801 CGTAATCACC CGAGTGTGAT CATCTGGTCG CTGGGGAATG AATCAGGCCA  
GCATTAGTGG GCTCACACTA GTAGACCAGC GACCCCTTAC TTAGTCCGGT

```
2851  CGGCGCTAAT CACGACGCGC TGTATCGCTG GATCAAATCT GTCGATCCTT
      GCCGCGATTA GTGCTGCGCG ACATAGCGAC CTAGTTTAGA CAGCTAGGAA
-----
2901  CCCGCCCCGGT GCAGTATGAA GGC GGCGGAG CCGACACCAC GGCCACCGAT
      GGGCGGGCCA CGTCATACTT CCGCCGCCTC GGCTGTGGTG CCGGTGGCTA
-----
2951  ATTATTTGCC CGATGTACGC GCGCGTGGAT GAAGACCAGC CCTTCCCGGC
      TAATAAACGG GCTACATGCG CGCGCACCTA CTTCTGGTCG GGAAGGGCCG
-----
3001  TGTGCCGAAA TGGTCCATCA AAAAATGGCT TTCGCTACCT GGAGAGACGC
      ACACGGCTTT ACCAGGTAGT TTTTACC GAAGCATGGA CCTCTCTGCG
-----
3051  GCCCGCTGAT CCTTTGCGAA TACGCCACG CGATGGGTAA CAGTCTTGGC
      CGGGCGACTA GGAAACGCTT ATGCGGGTGC GCTACCCATT GTCAGAACCG
-----
3101  GGTTCGCTA AATACTGGCA GGC GTTTCGT CAGTATCCCC GTTTACAGGG
      CCAAAGCGAT TTATGACCGT CCGCAAAGCA GTCATAGGGG CAAATGTCCC
-----
3151  CGGCTTCGTC TGGGACTGGG TGGATCAGTC GCTGATTAAA TATGATGAAA
      GCCGAAGCAG ACCCTGACCC ACCTAGTCAG CGACTAATTT ATACTACTTT
-----
3201  ACGGCAACCC GTGGTCGGCT TACGGCGGTG ATTTTGGCGA TACGCCGAAC
      TGCCGTTGGG CACCAGCCGA ATGCCGCCAC TAAAACCGCT ATGCGGCTTG
-----
3251  GATCGCCAGT TCTGTATGAA CGGTCTGGTC TTTGCCGACC GCACGCCGCA
      CTAGCGGTCA AGACATACTT GCCAGACCAG AAACGGCTGG CGTGCGCGCT
-----
3301  TCCAGCGCTG ACGGAAGCAA AACACCAGCA GCAGTTTTTC CAGTTCCGTT
      AGGTGCGCAG TGCCTTCGTT TTGTGGTCGT CGTCAAAAAG GTCAAGGCAA
-----
3351  TATCCGGGCA AACCATCGAA GTGACCAGCG AATACCTGTT CCGTCATAGC
      ATAGGCCCGT TTGGTAGCTT CACTGGTCGC TTATGGACAA GGCAGTATCG
-----
3401  GATAACGAGC TCCTGCACTG GATGGTGGCG CTGGATGGTA AGCCGCTGGC
      CTATTGCTCG AGGACGTGAC CTACCACCGC GACCTACCAT TCGGCGACCG
-----
3451  AAGCGGTGAA GTGCCTCTGG ATGTCGCTCC ACAAGGTAAA CAGTTGATTG
      TTCGCCACTT CACGGAGACC TACAGCGAGG TGTTCCATTT GTCAACTAAC
-----
3501  AACTGCCTGA ACTACCGCAG CCGGAGAGCG CCGGGCAACT CTGGCTCACA
      TTGACGGACT TGATGGCGTC GGCCTCTCGC GGCCCGTTGA GACCGAGTGT
-----
3551  GTACGCGTAG TGCAACCGAA CGCGACCGCA TGGTCAGAAG CCGGGCACAT
      CATGCGCATC ACGTTGGCTT GCGCTGGCGT ACCAGTCTTC GGCCCGTGTA
-----
3601  CAGCGCCTGG CAGCAGTGGC GTCTGGCGGA AAACCTCAGT GTGACGCTCC
      GTCGCGGACC GTCGTCACCG CAGACCGCCT TTTGGAGTCA CACTGCGAGG
-----
3651  CCGCCGCGTC CCACGCCATC CCGCATCTGA CCACCAGCGA AATGGATTTT
      GGCGGCGCAG GGTGCGGTAG GCGTAGACT GGTGGTCGCT TTACCTAAAA
-----
3701  TGCATCGAGC TGGGTAATAA GCGTTGGCAA TTTAACCGCC AGTCAGGCTT
      ACGTAGCTCG ACCCATATT CGCAACCGTT AAATTGGCGG TCAGTCCGAA
-----
3751  TCTTTCACAG ATGTGGATTG GCGATAAAAA ACAACTGCTG ACGCCGCTGC
      AGAAAGTGTC TACACCTAAC CGCTATTTT TGTTGACGAC TGCGGCGACG
-----
```

3801 GCGATCAGTT CACCCGTGTC GATAGATCTG AACAGAACT CATTTCGGAA  
CGCTAGTCAA GTGGGCACAG CTATCTAGAC TTGTCTTTGA GTAAAGGCTT

3851 GAAGACCTAG TCGACCATCA TCATCATCAT CACCGGTAAT AATAGGTAGA  
CTTCTGGATC AGCTGGTAGT AGTAGTAGTA GTGGCCATTA TTATCCATCT

3901 TAAGTGACTG ATTAGATGCA TTTCGACTAG ATCCCTCGAC CAATTCCGGT  
ATTCAGTAC TAATCTACGT AAAGCTGATC TAGGGAGCTG GTTAAGGCCA

3951 TATTTTCCAC CATATTGCCG TCTTTTGGCA ATGTGAGGGC CCGGAAACCT  
ATAAAAGGTG GTATAACGGC AGAAAACCGT TACACTCCCG GGCCTTTGGA

4001 GGCCCTGTCT TCTTGACGAG CATTCCTAGG GGTCTTTCCC CTCTCGCCAA  
CCGGGACAGA AGAACTGCTC GTAAGGATCC CCAGAAAGGG GAGAGCGGTT

4051 AGGAATGCAA GGTCTGTTGA ATGTCGTGAA GGAAGCAGTT CCTCTGGAAG  
TCCTTACGTT CCAGACAACT TACAGCACTT CCTTCGTCAA GGAGACCTTC

4101 CTTCTTGAAG ACAAACAACG TCTGTAGCGA CCCTTTGCAG GCAGCGGAAC  
GAAGAACTTC TGTTTGTGTC AGACATCGCT GGGAAACGTC CGTCGCCTTG

4151 CCCCCACCTG GCGACAGGTG CCTCTGCGGC CAAAAGCCAC GTGTATAAGA  
GGGGGTGGAC CGCTGTCCAC GGAGACGCCG GTTTTCGGTG CACATATTCT

4201 TACACCTGCA AAGGCGGCAC AACCCAGTG CCACGTTGTG AGTTGGATAG  
ATGTGGACGT TTCCGCCGTG TTGGGGTCAC GGTGCAACAC TCAACCTATC

4251 TTGTGAAAAG AGTCAAATGG CTCTCCTCAA GCGTATTCAA CAAGGGGCTG  
AACACCTTTC TCAGTTTACC GAGAGGAGTT CGCATAAGTT GTTCCCCGAC

4301 AAGGATGCCC AGAAGGTACC CCATTGTATG GGATCTGATC TGGGGCCTCG  
TTCTTACGGG TCTTCCATGG GGTAACATAC CCTAGACTAG ACCCCGGAGC

4351 GTGCACATGC TTTACATGTG TTTAGTCGAG GTTAAAAAAC GTCTAGGCCC  
CACGTGTACG AAATGTACAC AAATCAGCTC CAATTTTTTG CAGATCCGGG

4401 CCCGAACCAC GGGGACGTGG TTTTCCTTTG AAAAACACGA TGATAATACC  
GGGCTTGGTG CCCCTGCACC AAAAGGAAAC TTTTGTGCT ACTATTATGG

4451 ATGAAAAAGC CTGAACTCAC CGCGACGTCT GTCGAGAAGT TTCTGATCGA  
TACTTTTTTCG GACTTGAGTG GCGCTGCAGA CAGCTCTTCA AAGACTAGCT

4501 AAAGTTCGAC AGCGTCTCCG ACCTGATGCA GCTCTCGGAG GGCGAAGAAT  
TTTCAAGCTG TCGCAGAGGC TGGACTACGT CGAGAGCCTC CCGCTTCTTA

4551 CTCGTGCTTT CAGCTTCGAT GTAGGAGGGC GTGGATATGT CCTGCGGGTA  
GAGCACGAAA GTCGAAGCTA CATCTCCCG CACCTATACA GGACGCCCAT

4601 AATAGCTGCG CCGATGGTTT CTACAAAGAT CGTTATGTTT ATCGGCACTT  
TTATCGACGC GGCTACCAAA GATGTTTCTA GCAATACAAA TAGCCGTGAA

4651 TGCATCGGCC GCGCTCCCGA TTCCGGAAGT GCTTGACATT GGGGAATTTA  
ACGTAGCCGG CCGGAGGGCT AAGGCCTTCA CGAACTGTAA CCCCTTAAAT

4701 GCGAGAGCCT GACCTATTGC ATCTCCCGCC GTGCACAGGG TGTCACGTTG  
CGCTCTCGGA CTGGATAACG TAGAGGGCGG CACGTGTCCC ACAGTGCAAC

4751 CAAGACCTGC CTGAAACCGA ACTGCCCGCT GTTCTGCAGC CGGTCGCGGA  
GTTCTGGACG GACTTTGGCT TGACGGGCGA CAAGACGTCG GCCAGCGCCT

4801 GGCCATGGAT GCGATCGCTG CGGCCGATCT TAGCCAGACG AGCGGGTTCTG  
CCGGTACCTA CGCTAGCGAC GCCGGCTAGA ATCGGTCTGC TCGCCCAAGC

4851 GCCCATTTCGG ACCGCAAGGA ATCGGTCAAT AACTACATG GCGTGATTTTC  
CGGGTAAGCC TGGCGTTCCT TAGCCAGTTA TGTGATGTAC CGCACTAAAG

4901 ATATGCGCGA TTGCTGATCC CCATGTGTAT CACTGGCAAA CTGTGATGGA  
TATACGCGCT AACGACTAGG GGTACACATA GTGACCGTTT GACACTACCT

4951 CGACACCGTC AGTGCGTCCG TCGCGCAGGC TCTCGATGAG CTGATGCTTT  
GCTGTGGCAG TCACGCAGGC AGCGCGTCCG AGAGCTACTC GACTACGAAA

5001 GGGCCGAGGA CTGCCCCGAA GTCCGGCACC TCGTGCACGC GGATTTTCGGC  
CCGGGCTCCT GACGGGGCTT CAGGCCGTGG AGCACGTGCG CCTAAAGCCG

5051 TCCAACAATG TCCTGACGGA CAATGGCCGC ATAACAGCGG TCATTGACTG  
AGGTTGTTAC AGGACTGCCT GTTACCGGCG TATTGTCGCC AGTAACTGAC

5101 GAGCGAGGCG ATGTTTCGGG ATTCCCAATA CGAGGTCGCC AACATCTTCT  
CTCGCTCCGC TACAAGCCCC TAAGGGTTAT GCTCCAGCGG TTGTAGAAGA

5151 TCTGGAGGCC GTGGTTGGCT TGTATGGAGC AGCAGACGCG CTAATTCGAG  
AGACCTCCGG CACCAACCGA ACATACCTCG TCGTCTGCGC GATGAAGCTC

5201 CGGAGGCATC CGGAGCTTGC AGGATCGCCG CGGCTCCGGG CGTATATGCT  
GCCTCCGTAG GCCTCGAACG TCCTAGCGGC GCCGAGGCC GCATATACGA

5251 CCGCATTGGT CTTGACCAAC TCTATCAGAG CTTGGTTGAC GGCAATTTTCG  
GGCGTAACCA GAAGTGGTTG AGATAGTCTC GAACCAACTG CCGTTAAAGC

5301 ATGATGCAGC TTGGGCGCAG GGTGATGCG ACGCAATCGT CCGATCCGGA  
TACTACGTCTG AACCCGCGTC CCAGCTACGC TCGGTTAGCA GGCTAGGCCT

5351 GCCGGGACTG TCGGGCGTAC ACAAATCGCC CGCAGAAGCG CGGCCGTCTG  
CGGCCCTGAC AGCCCGCATG TGTTTAGCGG GCGTCTTCGC GCCGGCAGAC

5401 GACCGATGGC TGTGTAGAAG TACTCGCCGA TAGTGGAAC CGACGCCCCA  
CTGGCTACCG ACACATCTTC ATGAGCGGCT ATCACCTTTG GCTGCGGGGT

5451 GCACTCGTCC GAGGGCAAAG GAATAGAGTA GATGCCGACC GGGATCTATC  
CGTGAGCAGG CTCCCGTTTC CTTATCTCAT CTACGGCTGG CCCTAGATAG

5501 GATAAAATAA AAGATTTTAT TTAGTCTCCA GAAAAAGGGG GGAATGAAAG  
CTATTTTATT TTCTAAATA AATCAGAGGT CTTTTTCCCC CCTTACTTTC

5551 ACCCCACCTG TAGGTTTGGC AAGCTAGCTT AAGTAACGCC ATTTTGCAAG  
TGGGGTGGAC ATCCAAACCG TTCGATCGAA TTCATTGCGG TAAAACGTTT

5601 GCATGGAAAA ATACATAACT GAGAATAGAG AAGTTCAGAT CAAGGTCAGG  
CGTACCTTTT TATGTATTGA CTCTTATCTC TTCAAGTCTA GTTCCAGTCC

5651 AACAGATGGA ACAGCTGAAT ATGGGCCAAA CAGGATATCT GTGGTAAGCA  
TTGTCTACCT TGTCGACTTA TACCCGTTT GTCCTATAGA CACCATTTCGT

```

5701 GTTCCTGCCC CGGCTCAGGG CCAAGAACAG ATGGAACAGC TGAATATGGG
    CAAGGACGGG GCCGAGTCCC GGTTCCTTGC TACCTTGTCG ACTTATACCC
-----
5751 CCAAACAGGA TATCTGTGGT AAGCAGTTCC TGCCCCGGCT CAGGGCCAAG
    GGTTTGTCTT ATAGACACCA TTCGTCAAGG ACGGGGCCGA GTCCCCGGTC
-----
5801 AACAGATGGT CCCAGATGC GGTCCAGCCC TCAGCAGTTT CTAGAGAACC
    TTGTCTACCA GGGGTCTACG CCAGGTCGGG AGTCGTCAAA GATCTCTTGG
-----
5851 ATCAGATGTT TCCAGGGTGC CCCAAGGACC TGAAATGACC CTGTGCCTTA
    TAGTCTACAA AGGTCCCACG GGGTTCCTGG ACTTTACTGG GACACGGAAT
-----
5901 TTTGAACTAA CCAATCAGTT CGCTTCTCGC TTCTGTTCGC GCGCTTCTGC
    AAACCTTGATT GGTTAGTCAA GCGAAGAGCG AAGACAAGCG CGCGAAGACG
-----
5951 TCCCCGAGCT CAATAAAAGA GCCCACAACC CCTCACTCGG GCGGCCAGTC
    AGGGGCTCGA GTTATTTTCT CGGGTGTGGG GGAGTGAGCC CCGCGGTCAG
-----
6001 CTCCGATTGA CTGAGTCGCC CGGGTACCCG TGTATCCAAT AAACCCTCTT
    GAGGCTAACT GACTCAGCGG GCCCATGGGC ACATAGGTTA TTTGGGAGAA
-----
6051 GCAGTTGCAT CCGACTTGTG GTCTCGCTGT TCCTTGGGAG GGTCTCCTCT
    CGTCAACGTA GGTGGAACAC CAGAGCGACA AGGAACCCTC CCAGAGGAGA
-----
6101 GAGTGATTGA CTACCCGTCA GCGGGGGTCT TTCATTCTAT CAGCATGTAT
    CTCCTAACT GATGGGCAGT CGCCCCCAGA AAGTAAGTAC GTCGTACATA
-----
6151 CAAAATTAAT TTGGTTTTTT TTCTTAAGTA TTTACATTAA ATGGCCATAG
    GTTTTAATTA AACCAAAAAA AAGAATTCAT AAATGTAATT TACCGGTATC
-----
6201 TTGCATTAAT GAATCGGCCA ACGCGCGGGG AGAGGCGGTT TGCGTATTGG
    AACGTAATTA CTTAGCCGGT TGCGCGCCCC TCTCCGCCAA ACGCATAACC
-----
6251 CGCTCTTCCG CTTCTCTGCT CACTGACTCG CTGCGCTCGG TCGTTCGGCT
    GCGAGAAGGC GAAGGAGCGA GTGACTGAGC GACGCGAGCC AGCAAGCCGA
-----
6301 GCGGCGAGCG GTATCAGCTC ACTCAAAGGC GGTAATACGG TTATCCACAG
    CGCCGCTCGC CATAGTCGAG TGAGTTTCCG CCATTATGCC AATAGGTGTC
-----
6351 AATCAGGGGA TAACGCAGGA AAGAACATGT GAGCAAAAGG CCAGCAAAAG
    TTAGTCCCCT ATTGCGTCCT TTCTTGTAACA CTCGTTTTCC GGTGTTTTTC
-----
6401 GCCAGGAACC GTAAAAAGGC CGCGTTGCTG GCGTTTTTCC ATAGGCTCCG
    CGGTCCTTGG CATTTTTCCG GCGCAACGAC CGCAAAAAGG TATCCGAGGC
-----
6451 CCCCCCTGAC GAGCATCACA AAAATCGACG CTCAAGTCAG AGGTGGCGAA
    GGGGGGACTG CTCGTAGTGT TTTTAGCTGC GAGTTCAGTC TCCACCGCTT
-----
6501 ACCCGACAGG ACTATAAAGA TACCAGGCGT TTCCCCCTGG AAGCTCCCTC
    TGGGCTGTCC TGATATTTCT ATGGTCCGCA AAGGGGGACC TTCGAGGGAG
-----
6551 GTGCGCTCTC CTGTTCGAC CCTGCCGCTT ACCGGATACC TGTCCGCCTT
    CACGCGAGAG GACAAGGCTG GGACGGCGAA TGGCCTATGG ACAGGCGGAA
-----
6601 TCTCCCTTCG GGAAGCGTGG CGCTTTCTCA TAGCTCACGC TGTAGGTATC
    AGAGGGAAGC CCTTCGCACC GCGAAAGAGT ATCGAGTGCG ACATCCATAG
-----

```

```

6651 TCAGTTCGGT GTAGGTCGTT CGCTCCAAGC TGGGCTGTGT GCACGAACCC
    AGTCAAGCCA CATCCAGCAA GCGAGGTTTCG ACCCGACACA CGTGCTTGGG
-----
6701 CCCGTTTCAGC CCGACCGCTG CGCCTTATCC GGTAACATATC GTCTTGAGTC
    GGGCAAGTCG GGCTGGCGAC GCGGAATAGG CCATTGATAG CAGAAGTCAG
-----
6751 CAACCCGGTA AGACACGACT TATCGCCACT GGCAGCAGCC ACTGGTAACA
    GTTGGGCCAT TCTGTGCTGA ATAGCGGTGA CCGTCGTCGG TGACCATTGT
-----
6801 GGATTAGCAG AGCGAGGTAT GTAGGCGGTG CTACAGAGTT CTTGAAGTGG
    CCTAATCGTC TCGCTCCATA CATCCGCCAC GATGTCTCAA GAACTTCACC
-----
6851 TGGCCTAACT ACGGCTACAC TAGAAGAACA GTATTTGGTA TCTGCGCTCT
    ACCGGATTGA TGCCGATGTG ATCTTCTTGT CATAAACCAT AGACGCGAGA
-----
6901 GCTGAAGCCA GTTACCTTCG GAAAAAGAGT TGGTAGCTCT TGATCCGGCA
    CGACTTCGGT CAATGGAAGC CTTTTTCTCA ACCATCGAGA ACTAGGCCGT
-----
6951 AACAAACCAC CGCTGGTAGC GGTGGTTTTT TTGTTTGCAA GCAGCAGATT
    TTGTTTGGTG GCGACCATCG CCACCAAAAA AACAAACGTT CGTCGTCTAA
-----
7001 ACGCGCAGAA AAAAAGGATC TCAAGAAGAT CCTTTGATCT TTTCTACGGG
    TGC GCGTCTT TTTTTCCTAG AGTCTTCTA GGAACTAGA AAAGATGCCC
-----
7051 GTCTGACGCT CAGTGAACG AAAACTCACG TTAAGGGATT TTGGTCATGA
    CAGACTGCGA GTCACCTTGC TTTTGAGTGC AATTCCCTAA AACCAGTACT
-----
7101 GATTATCAAA AAGGATCTTC ACCTAGATCC TTTTAAATTA AAAATGAAGT
    CTAATAGTTT TTCCTAGAAG TGGATCTAGG AAAATTTAAT TTTTACTTCA
-----
7151 TTGCGGCCGC AAATCAATCT AAAGTATATA TGAGTAAACT TGGTCTGACA
    AACGCCGGCG TTTAGTTAGA TTTCATATAT ACTCATTTGA ACCAGACTGT
-----
7201 GTTACCAATG CTTAATCAGT GAGGCACCTA TCTCAGCGAT CTGTCTATTT
    CAATGGTTAC GAATTAGTCA CTCCGTGGAT AGAGTCGCTA GACAGATAAA
-----
7251 CGTTCATCCA TAGTTGCTG ACTCCCGTC GTGTAGATAA CTACGATACG
    GCAAGTAGGT ATCAACGGAC TGAGGGGCAG CACATCTATT GATGCTATGC
-----
7301 GGAGGGCTTA CCATCTGGCC CCAGTGCTGC AATGATACCG CGAGACCCAC
    CCTCCCGAAT GGTAGACCGG GGTCACGACG TTACTATGGC GCTCTGGGTG
-----
7351 GCTCACC GGC TCCAGATTTA TCAGCAATAA ACCAGCCAGC CGGAAGGGCC
    CGAGTG GCGG AGGTCTAAAT AGTCGTTATT TGGTCGGTCG GCCTTCCCGG
-----
7401 GAGCGCAGAA GTGGTCCTGC AACTTTATCC GCCTCCATCC AGTCTATTAA
    CTCGCGTCTT CACCAGGACG TTGAAATAGG CGGAGGTAGG TCAGATAATT
-----
7451 TTGTTGCCGG GAAGCTAGAG TAAGTAGTTC GCCAGTTAAT AGTTTGCGCA
    AACAACGGCC CTTCGATCTC ATTCATCAAG CGGTCAATTA TCAAACGCGT
-----
7501 ACGTTGTTGC CATTGCTACA GGCATCGTGG TGTCACGCTC GTCGTTTGGT
    TGCAACAACG GTAACGATGT CCGTAGCACC ACAGTGCAG CAGCAAACCA
-----
7551 ATGGCTTCAT TCAGCTCCGG TTCCCAACGA TCAAGGCGAG TTACATGATC
    TACCGAAGTA AGTCGAGGCC AAGGGTTGCT AGTTCCGCTC AATGTACTAG
-----

```



7601 CCCCATGTTG TGCAAAAAAG CGGTTAGCTC CTTGGGTCCT CCGATCGTTG  
GGGGTACAAC ACGTTTTTTC GCCAATCGAG GAAGCCAGGA GGCTAGCAAC

7651 TCAGAAGTAA GTTGGCCGCA GTGTTATCAC TCATGGTTAT GGCAGCACTG  
AGTCTTCATT CAACCGGCGT CACAATAGTG AGTACCAATA CCGTCGTGAC

7701 CATAATTCTC TTAAGTGCAT GCCATCCGTA AGATGCTTTT CTGTGACTGG  
GTATTAAGAG AATGACAGTA CGGTAGGCAT TCTACGAAAA GACACTGACC

7751 TGAGTACTCA ACCAAGTCAT TCTGAGAATA GTGTATGCGG CGACCGAGTT  
ACTCATGAGT TGGTTCAGTA AGACTCTTAT CACATACGCC GCTGGCTCAA

7801 GCTCTTGCCC GCGGTCAATA CGGGATAATA CCGCGCCACA TAGCAGAACT  
CGAGAACGGG CCGCAGTTAT GCCCTATTAT GCGCGGGTGT ATCGTCTTGA

7851 TTAAGAGTGC TCATCATTGG AAAACGTTCT TCGGGGCGAA AACTCTCAAG  
AATTTTCACG AGTAGTAACC TTTTGCAAGA AGCCCCGCTT TTGAGAGTTC

7901 GATCTTACCG CTGTTGAGAT CCAGTTCGAT GTAACCCACT CGTGCACCCA  
CTAGAATGGC GACAACTCTA GGTCAAGCTA CATTGGGTGA GCACGTGGGT

7951 ACTGATCTTC AGCATCTTTT ACTTTCACCA GCGTTTCTGG GTGAGCAAAA  
TGACTAGAAG TCGTAGAAAA TGAAAGTGGT CGCAAAGACC CACTCGTTTT

8001 ACAGGAAGGC AAAATGCCGC AAAAAAGGGA ATAAGGGCGA CACGGAAATG  
TGCCTTCCG TTTTACGGCG TTTTTCCTT TATTCCCGCT GTGCCTTTAC

8051 TTGAATACTC ATACTCTTCC TTTTCAATA TTATTGAAGC ATTTATCAGG  
AACTTATGAG TATGAGAAGG AAAAAGTTAT AATAACTTCG TAAATAGTCC

8101 GTTATTGTCT CATGAGCGGA TACATATTTG AATGTATTTA GAAAAATAAA  
CAATAACAGA GTACTCGCCT ATGTATAAAC TTACATAAAT CTTTTTATTT

8151 CAAATAGGGG TTCCGCGCAC ATTTT  
GTTTATCCCC AAGGCGCGTG TAAAG

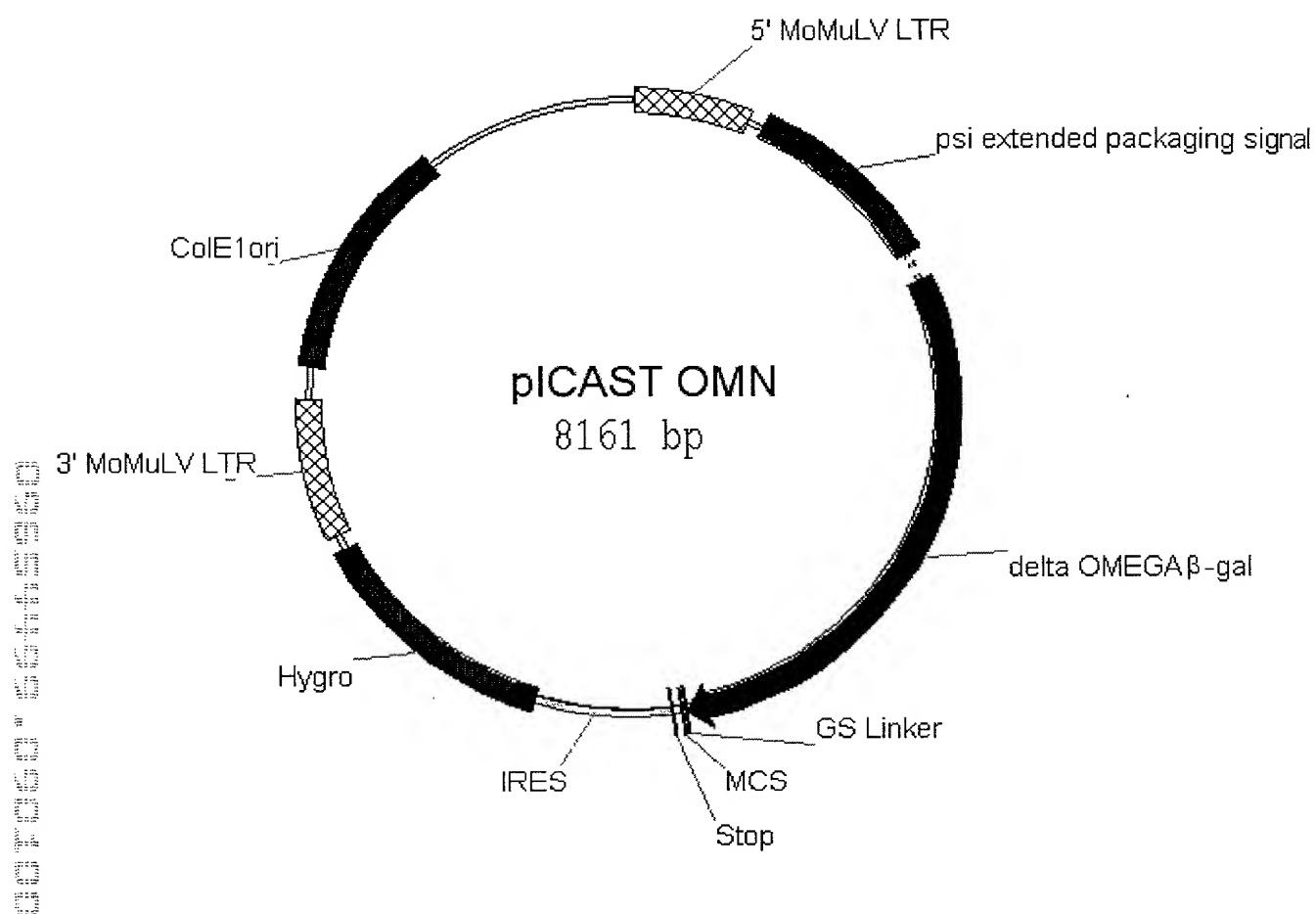


Figure 13A

```

1  CTGCAGCCTG AATATGGGCC AAACAGGATA TCTGTGGTAA GCAGTTCCTG
   GACGTCGGAC TTATACCCGG TTTGTCCTAT AGACACCATT CGTCAAGGAC
-----
51  CCCC GGCTCA GGGCCAAGAA CAGATGGAAC AGCTGAATAT GGGCCAAACA
   GGGGCCGAGT CCCGGTTCTT GTCTACCTTG TCGACTTATA CCCGGTTTGT
-----
101 GGATATCTGT GGTAAGCAGT TCCTGCCCCG GCTCAGGGCC AAGAACAGAT
   CCTATAGACA CCATTCGTCA AGGACGGGGC CGAGTCCCGG TTCTTGTCTA
-----
151 GGTCCCCAGA TGC GGTCAG CCCTCAGCAG TTTCTAGAGA ACCATCAGAT
   CCAGGGGTCT ACGCCAGGTC GGGAGTCGTC AAAGATCTCT TGGTAGTCTA
-----
201 GTTCCAGGG TGGCCCAAGG ACCTGAAATG ACCCTGTGCC TTATTTGAAC
   CAAAGGTCCC ACGGGGTTC TGGACTTTAC TGGGACACGG AATAAACTTG
-----
251 TAACCAATCA GTTCGCTTCT CGCTTCTGTT CGCGCGCTTC TGCTCCCCGA
   ATTGGTTAGT CAAGCGAAGA GCGAAGACAA GCGCGCGAAG ACGAGGGGCT
-----
301 GCTCAATAAA AGAGCCCACA ACCCTCACT CGGGGCGCCA GTCCTCCGAT
   CGAGTTATTT TCTCGGGTGT TGGGGAGTGA GCGGCGCGGT CAGGAGGCTA
-----
351 TGA CTGAGTC GCGCGGGTAC CCGTGTATCC AATAAACCCCT CTG CAGTTG
   ACTGACTCAG CCGGGCCCATG GGCACATAGG TTATTTGGGA GAACGTCAAC
-----
401 CATCCGACTT GTGGTCTCGC TGTTCTTGG GAGGGTCTCC TCTGAGTGAT
   GTAGGCTGAA CACCAGAGCG ACAAGGAACC CTCCAGAGG AACTCACTA
-----
451 TGA CTACCCG TCAGCGGGGG TCTTTCATTT GGGGGCTCGT CCGGGATCGG
   ACTGATGGGC AGTCGCCCCC AGAAAGTAAA CCGCGAGCA GGCCCTAGCC
-----
501 GAGACCCCTG CCCAGGGACC ACCGACCCAC CACCGGGAGG CAAGCTGGCC
   CTCTGGGGAC GGGTCCCTGG TGGCTGGGTG GTGGCCCTCC GTTCGACCGG
-----
551 AGCAACTTAT CTGTGTCTGT CCGATTGTCT AGTGTCTATG ACTGATTTTA
   TCGTTGAATA GACACAGACA GGCTAACAGA TCACAGATAC TGA TAAAT
-----
601 TGCGCCTGCG TCGGTACTAG TTAGCTAACT AGCTCTGTAT CTGGCGGACC
   ACGCGGACGC AGCCATGATC AATCGATTGA TCGAGACATA GACCGCCTGG
-----
651 CGTGGTGGA CTGACGAGTT CTGAACACCC GGCCGCAACC CTGGGAGACG
   GCACCACCTT GACTGCTCAA GACTTGTGGG CCGGCGTTGG GACCTCTGC
-----
701 TCCCAGGGAC TTTGGGGGCC GTTTTGTGG CCCGACCTGA GGAAGGGAGT
   AGGGTCCCTG AAACCCCGG CAAAAACACC GGGCTGGACT CCTTCCCTCA
-----
751 CGATGTGGAA TCCGACCCCG TCAGGATATG TGGTTCTGGT AGGAGACGAG
   GCTACACCTT AGGCTGGGGC AGTCCTATAC ACCAAGACCA TCCTCTGCTC
-----
801 AACCTAAAAC AGTTCCCGCC TCCGTCTGAA TTTTGTCTT CGGTTTGGAA
   TTGGATTTTG TCAAGGGCGG AGGCAGACTT AAAAAAGAAA GCCAAACCTT
-----
851 CCGAAGCCGC GCGTCTTGTC TGCTGCAGCA TCGTCTGTG TTGTCTCTGT
   GGCTTCGGCG CGCAGAACAG ACGACGTCGT AGCAAGACAC AACAGAGACA
-----
901 CTGACTGTGT TTCTGTATTT GTCTGAAAAT TAGGGCCAGA CTGTTACCAC
   GACTGACACA AAGACATAAA CAGACTTTTA ATCCCGGTCT GACAATGGTG
-----

```

FIGURE 13B

```

951  TCCCTTAAGT TTGACCTTAG GTAAC TGGAA AGATGTCGAG CGGCTCGCTC
     AGGGAATTCA AACTGGAATC CATTGACCTT TCTACAGCTC GCCGAGCGAG
-----
1001  ACAACCAAGTC GGTAGATGTC AAGAAGAGAC GTTGGGTAC CTTCTGCTCT
     TGTGTCGAG CCATCTACAG TTCTTCTCTG CAACCAATG GAAGACGAGA
-----
1051  GCAGAATGGC CAACCTTTAA CGTCGGATGG CCGCGAGACG GCACCTTTAA
     CGTCTTACCG GTTGGAATTT GCAGCCTACC GCGCTCTGC CGTGGAATTT
-----
1101  CCGAGACCTC ATCACCAGG TTAAGATCAA GGTCTTTTCA CCTGGCCCCG
     GGCTCTGGAG TAGTGGGTCC AATTCTAGTT CCAGAAAAGT GGACCGGGCG
-----
1151  ATGGACACCC AGACCAGGTC CCCTACATCG TGACCTGGGA AGCCTTGGCT
     TACCTGTGGG TCTGGTCCAG GGGATGTAGC ACTGGACCTT TCGGAACCGA
-----
1201  TTTGACCCCC CTCCCTGGGT CAAGCCCTTT GTACACCCTA AGCCTCCGCC
     AAACCTGGGG GAGGGACCCA GTTCGGGAAA CATGTGGGAT TCGGAGGCGG
-----
1251  TCCTCTTCTT CCATCCGCCC CGTCTCTCCC CTTTGAACCT CCTCGTTTCA
     AGGAGAAGGA GGTAGGCGGG GCAGAGAGGG GGAACCTTGA GGAGCAAGCT
-----
1301  CCCC GCCTCG ATCCTCCCTT TATCCAGCCC TCACTCCTTC TCTAGGCGCC
     GGGGCGGAGC TAGGAGGGAA ATAGGTCGGG AGTGAGGAAG AGATCCGCGG
-----
1351  GGCCGCTCTA GCCCATTAAAT ACGACTCACT ATAGGGCGAT TCGAACACCA
     CCGGCGAGAT CGGGTAATTA TGCTGAGTGA TATCCCGCTA AGCTTGTGGT
-----
1401  TGCACCATCA TCATCATCAC GTCGACGAAC AGAAACTCAT TTCCGAAGAA
     ACGTGGTAGT AGTAGTAGTG CAGCTGCTTG TCTTTGAGTA AAGGCTTCTT
-----
1451  GACCTACTCG AGATGGGCGT GATTACGGAT TCACTGGCCG TCGTTTACA
     CTGGATGAGC TCTACCCGCA CTAATGCCTA AGTGACCGGC AGCAAAATGT
-----
1501  ACGTCGTGAC TGGGAAAACC CTGGCGTTAC CCAACTTAAT CGCCTTGCAG
     TGCAGCACTG ACCCTTTTGG GACCGCAATG GGTGAATTA GCGGAACGTC
-----
1551  CACATCCCCC TTTGCCAGC TGGCGTAATA GCGAAGAGGC CCGCACCGAT
     GTGTAGGGGG AAAGCGGTCG ACCGCATTAT CGCTTCTCCG GCGTGGCTA
-----
1601  CGCCCTTCCC AACAGTTACG CAGCCTGAAT GGCGAATGGC GCTTTGCCTG
     GCGGGAAGGG TTGTCAATGC GTCGGACTTA CCGCTTACCG CGAAACGGAC
-----
1651  GTTTCGGGCA CCAGAAGCGG TGCCGGAAG CTGGCTGGAG TGCGATCTTC
     CAAAGGCCGT GGTCTTCGCC ACGGCCTTTC GACCGACCTC ACGCTAGAAG
-----
1701  CTGAGGCCGA TACTGTCTGC GTCCCTCAA ACTGGCAGAT GCACGGTTAC
     GACTCCGGCT ATGACAGCAG CAGGGGAGTT TGACCGTCTA CGTGCCAATG
-----
1751  GATGCGCCCA TCTACACCAA CGTGACCTAT CCCATTACGG TCAATCCGCC
     CTACGCGGGT AGATGTGGTT GCACTGGATA GGGTAATGCC AGTTAGGCGG
-----
1801  GTTTGTTCCC ACGGAGAATC CGACGGGTTG TTA CTGCTC ACATTTAATG
     CAAACAAGGG TGCTCTTAG GCTGCCAAC AATGAGCGAG TGTAATTAC
-----
1851  TTGATGAAAG CTGGCTACAG GAAGGCCAGA CGCGAATTAT TTTTGATGGC
     AACTACTTTC GACCGATGTC CTTCCGGTCT GCGCTTAATA AAAACTACCG
-----

```

```

1901 GTTAACTCGG CGTTTCATCT GTGGTGCAAC GGGCGCTGGG TCGGTTACGG
    CAATTGAGCC GCAAAGTAGA CACCACGTTG CCCGCGACCC AGCCAATGCC
-----
1951 CCAGGACAGT CGTTTGCCGT CTGAATTTGA CCTGAGCGCA TTTTACGCG
    GGTCTGTCA GCAAACGGCA GACTTAACT GGA CTGCGT AAAAATGCGC
-----
2001 CCGGAGAAAA CCGCCTCGCG GTGATGGTGC TGCCTGGAG TGACGGCAGT
    GGCCTCTTTT GCGGAGCGC CACTACCACG ACGCGACCTC ACTGCCGTCA
-----
2051 TATCTGGAAG ATCAGGATAT GTGGCGGATG AGCGGCATTT TCCGTGACGT
    ATAGACCTTC TAGTCCTATA CACCGCCTAC TCGCCGTAAG AGGCACTGCA
-----
2101 CTCGTTGCTG CATAAACCGA CTACACAAAT CAGCGATTTT CATGTTGCCA
    GAGCAACGAC GTATTGGGCT GATGTGTTTA GTCGCTAAAG GTACAACGGT
-----
2151 CTCGCTTTAA TGATGATTTT AGCCGCGCTG TACTGGAGGC TGAAGTTCAG
    GAGCGAAATT ACTACTAAAG TCGGCGCGAC ATGACCTCCG ACTTCAAGTC
-----
2201 ATGTGCGGCG AGTTGCGTGA CTACCTACGG GTAACAGTTT CTTTATGGCA
    TACACGCCGC TCAACGCACT GATGGATGCC CATTGTCAAA GAAATACCGT
-----
2251 GGGTGAAACG CAGGTCGCCA GCGGCACCGC GCCTTTCGGC GGTGAAATTA
    CCCACTTTGC GTCCAGCGGT CGCCGTGGCG CGGAAAGCCG CCACTTTAAT
-----
2301 TCGATGAGCG TGGTGGTTAT GCCGATCGCG TCACACTACG TCTGAACGTC
    AGCTACTCGC ACCACCAATA CGGCTAGCGC AGTGTGATGC AGACTTGCAAG
-----
2351 GAAAACCCGA AACTGTGGAG CGCCGAAATC CCGAATCTCT ATCGTGCGGT
    CTTTTGGGCT TTGACACCTC GCGGCTTTAG GGCTTAGAGA TAGCACGCCA
-----
2401 GGTGAACTG CACACCGCCG ACGGCACGCT GATTGAAGCA GAAGCCTGCG
    CCAACTTGAC GTGTGGCGGC TGCCGTGCGA CTAACCTCGT CTTCGGACGC
-----
2451 ATGTCGGTTT CCGCGAGGTG CGGATTGAAA ATGGTCTGCT GCTGCTGAAC
    TACAGCCAAA GCGGCTCCAC GCCTAACTTT TACCAGACGA CGACGACTTG
-----
2501 GGCAAGCCGT TGCTGATTCG AGGCGTTAAC CGTCACGAGC ATCATCCTCT
    CCGTTCGGCA ACGACTAAGC TCCGCAATTG GCAGTGCTCG TAGTAGGAGA
-----
2551 GCATGGTCAG GTCATGGATG AGCAGACGAT GGTGCAGGAT ATCCTGCTGA
    CGTACCAGTC CAGTACCTAC TCGTCTGCTA CCACGTCCTA TAGGACGACT
-----
2601 TGAAGCAGAA CAACTTTAAC GCCGTGCGCT GTTCGCATTA TCCGAACCAT
    ACTTCGTCTT GTTGAAATTG CGGCACGCGA CAAGCGTAAT AGGCTTGGTGTA
-----
2651 CCGCTGTGGT ACACGCTGTG CGACCGCTAC GGCCTGTATG TGGTGGATGA
    GGCGACACCA TGTGCGACAC GTGGCGATG CCGGACATAC ACCACCTACT
-----
2701 AGCCAATATT GAAACCCACG GCATGGTGCC AATGAATCGT CTGACCGATG
    TCGGTTATAA CTTTGGGTGC CGTACCACGG TTACTTAGCA GACTGGCTAC
-----
2751 ATCCGCGCTG GCTACCGGCG ATGAGCGAAC GCGTAACGCG AATGGTGCAG
    TAGGCGCGAC CGATGGCCGC TACTCGCTTG CGCATTGCGC TTACCACGTC
-----
2801 CGCGATCGTA ATCACCCGAG TGTGATCATC TGGTCGCTGG GGAATGAATC
    GCGCTAGCAT TAGTGGGCTC ACACCTAGTAG ACCAGCGACC CCTTACTTAG
-----

```

```
2851 AGGCCACGGC GCTAATCACG ACGCGCTGTA TCGCTGGATC AAATCTGTCG
TCCGGTGCCG CGATTAGTGC TGC GCGACAT AGCGACCTAG TTTAGACAGC
-----
2901 ATCCTTCCCG CCCGGTGCAG TATGAAGGCG GCGGAGCCGA CACCACGGCC
TAGGAAGGGC GGGCCACGTC ATACTTCCGC CGCCTCGGCT GTGGTGCCGG
-----
2951 ACCGATATTA TTTGCCCGAT GTACGCGCGC GTGGATGAAG ACCAGCCCTT
TGGCTATAAT AAACGGGCTA CATGCGCGCG CACCTACTTC TGGTCGGGAA
-----
3001 CCCGGCTGTG CCGAAATGGT CCATCAAAAA ATGGCTTTTCG CTACCTGGAG
GGGCCGACAC GGCTTTACCA GGTAGTTTTT TACCGAAAGC GATGGACCTC
-----
3051 AGACGCGCCC GCTGATCCTT TGCGAATACG CCCACGCGAT GGGTAACAGT
TCTGCGCGGG CGACTAGGAA ACGCTTATGC GGGTGCGCTA CCCATTGTCA
-----
3101 CTTGGCGGTT TCGCTAAATA CTGGCAGGCG TTTCGTCAGT ATCCCCGTTT
GAACCGCCAA AGCGATTAT GACCGTCCGC AAAGCAGTCA TAGGGGCAA
-----
3151 ACAGGGCGGC TTCGTCTGGG ACTGGGTGGA TCAGTCGCTG ATTAAATATG
TGTCGCCCGG AAGCAGACCC TGACCCACCT AGTCAGCGAC TAATTTATAC
-----
3201 ATGAAAACGG CAACCCGTGG TCGGCTTACG GCGGTGATTT TGGCGATACG
TACTTTTGCC GTTGGGCACC AGCCGAATGC CGCCACTAAA ACCGCTATGC
-----
3251 CCGAACGATC GCCAGTTCTG TATGAACGGT CTGGTCTTTG CCGACCGCAC
GGCTTGCTAG CGGTCAAGAC ATACTTGCCA GACCAGAAAC GGCTGGCGTG
-----
3301 GCCGCATCCA GCGCTGACGG AAGCAAAACA CCAGCAGCAG TTTTCCAGT
CGGCGTAGGT CGCGACTGCC TTCGTTTTGT GGTGCTGCTC AAAAAGGTCA
-----
3351 TCCGTTTATC CGGGCAAACC ATCGAAGTGA CCAGCGAATA CCTGTTCCGT
AGGCAAATAG GCCCGTTTGG TAGCTTCACT GGTGCTTAT GGACAAGGCA
-----
3401 CATAGCGATA ACGAGCTCCT GCACTGGATG GTGGCGCTGG ATGGTAAGCC
GTATCGCTAT TGCTCGAGGA CGTGACCTAC CACCGCGACC TACCATTTCG
-----
3451 GCTGGCAAGC GGTGAAGTGC CTCTGGATGT CGCTCCACAA GGTAACAGT
CGACCGTTCG CCACTTCACG GAGACCTACA GCGAGGTGTT CCATTGTCA
-----
3501 TGATTGAACT GCCTGAACTA CCGCAGCCGG AGAGCGCCGG GCAACTCTGG
ACTAACTTGA CGGACTTGAT GGCGTCGGCC TCTCGCGGCC CGTTGAGACC
-----
3551 CTCACAGTAC GCGTAGTGCA ACCGAACGCG ACCGCATGGT CAGAAGCCGG
GAGTGTCTAT CGCATCACGT TGGCTTGCGC TGGCGTACCA GTCTTCGGCC
-----
3601 GCACATCAGC GCCTGGCAGC AGTGGCGTCT GCGGGAAC CTCAGTGTGA
CGTGTAGTCG CGGACCGTCG TCACCGCAGA CCGCCTTTT GAGTCACACT
-----
3651 CGCTCCCCGC CGCGTCCCAC GCCATCCCCG ATCTGACCAC CAGCGAAATG
GCGAGGGGCG GCGCAGGGTG CGGTAGGGCG TAGACTGGTG GTCGCTTTAC
-----
3701 GATTTTTGCA TCGAGCTGGG TAATAAGCGT TGGCAATTA ACCGCCAGTC
CTAAAAACGT AGCTCGACCC ATTATTCGCA ACCGTTAAAT TGGCGGTCAG
-----
3751 AGGCTTTCTT TCACAGATGT GGATTGGCGA TAAAAACAA CTGCTGACGC
TCCGAAAGAA AGTGCTACA CCTAACGCT ATTTTTTGT GACGACTGCG
-----
```

```

3801  CGCTGCGCGA TCAGTTCACC CGTGTGCGATA GATCTGGAGG TGGTGGCAGC
      GCGACGCGCT AGTCAAGTGG GCACAGCTAT CTAGACCTCC ACCACCGTCG
-----
3851  AGGCCTTGGC GCGCCGGATC CTTAATTAAC AATTGACCGG TAATAATAGG
      TCCGGAACCG CGCGGCCTAG GAATTAATTG TTAAGTGGCC ATTATTATCC
-----
3901  TAGATAAGTG ACTGATTAGA TGCATTTCTGA CTAGATCCCT CGACCAATTC
      ATCTATTACAC TGAATAATCT ACGTAAAGCT GATCTAGGGA GCTGGTTAAG
-----
3951  CGGTTATTTT CCACCATATT GCCGTCTTTT GGCAATGTGA GGGCCCGGAA
      GCCAATAAAA GGTGTTATTA CGGCAGAAAA CCGTTACACT CCCGGGCCTT
-----
4001  ACCTGGCCCT GTCTTCTTGA CGAGCATTCG TAGGGGTCTT TCCCCTCTCG
      TGGACCGGGA CAGAAGAACT GCTCGTAAGG ATCCCCAGAA AGGGGAGAGC
-----
4051  CCAAAGGAAT GCAAGGTCTG TTGAATGTCG TGAAGGAAGC AGTTCCTCTG
      GGTTTCCTTA CGTTCAGAC AACTTACAGC ACTTCCTTCG TCAAGGAGAC
-----
4101  GAAGCTTCTT GAAGACAAAC AACGTCTGTA GCGACCCTTT GCAGGCAGCG
      CTTCAAGAA CTTCTGTTTG TTGCAGACAT CGCTGGGAAA CGTCCGTCGC
-----
4151  GAACCCCCCA CCTGGCGACA GGTGCCTCTG CGGCCAAAAG CCACGTGTAT
      CTTGGGGGGT GGACCGTGT CCACGGAGAC GCCGGTTTTT GGTGCACATA
-----
4201  AAGATACACC TGCAAAGGCG GCACAACCCC AGTGCCACGT TGTGAGTTGG
      TTCTATGTGG ACGTTTCCGC CGTGTGGGG TCACGGTGCA AACTCAACC
-----
4251  ATAGTTGTGG AAAGAGTCAA ATGGCTCTCC TCAAGCGTAT TCAACAAGGG
      TATCAACACC TTTCTCAGTT TACCGAGAGG AGTTCGCATA AGTTGTTCCC
-----
4301  GCTGAAGGAT GCCCAGAAGG TACCCCATTTG TATGGGATCT GATCTGGGGC
      CGACTTCCTA CGGGTCTTCC ATGGGGTAAC ATACCCTAGA CTAGACCCCG
-----
4351  CTCGGTGCAC ATGCTTTACA TGTGTTTAGT CGAGGTTAAA AAACGTCTAG
      GAGCCACGTG TACGAAATGT ACACAAATCA GCTCCAATTT TTTGCAGATC
-----
4401  GCCCCCGGAA CCACGGGGAC GTGGTTTTCC TTGAAAAAC ACGATGATAA
      CGGGGGGCTT GGTGCCCCTG CACCAAAAGG AAACTTTTTG TGCTACTATT
-----
4451  TACCATGAAA AAGCCTGAAC TCACGCGGAC GTCTGTGCGAG AAGTTTCTGA
      ATGGTACTTT TTCGGAATTG AGTGGCGCTG CAGACAGCTC TTCAAAGACT
-----
4501  TCGAAAAGTT CGACAGCGTC TCCGACCTGA TGCAGCTCTC GGAGGGCGAA
      AGCTTTTCAA GCTGTGCGAG AGGCTGGACT ACGTCGAGAG CCTCCCGCTT
-----
4551  GAATCTCGTG CTTTCAGCTT CGATGTAGGA GGGCGTGGAT ATGTCCTGCG
      CTTAGAGCAC GAAAGTCGAA GCTACATCCT CCCGCACCTA TACAGGACGC
-----
4601  GGTAATAGC TGCGCCGATG GTTTCTACAA AGATCGTTAT GTTTATCGGC
      CCATTTATCG ACGCGGCTAC CAAAGATGTT TCTAGCAATA CAAATAGCCG
-----
4651  ACTTTGCATC GGCCGCGCTC CCGATTCCGG AAGTGCTTGA CATTGGGGAA
      TGAAACGTAG CCGGCGCGAG GGCTAAGGCC TTCACGAACT GTAACCCCTT
-----
4701  TTTAGCGAGA GCCTGACCTA TTGCATCTCC CGCCGTGCAC AGGGTGTAC
      AAATCGCTCT CGGACTGGAT AACGTAGAGG GCGGCACGTG TCCCACAGTG
-----

```

```

4751 GTTGCAAGAC CTGCCTGAAA CCGAACTGCC CGCTGTTCTG CAGCCGGTCCG
      CAACGTTCTG GACGGACTTT GGCTTGACGG GCGACAAGAC GTCGGCCAGC
-----
4801 CGGAGGCCAT GGATGCGATC GCTGCGGCCG ATCTTAGCCA GACGAGCGGG
      GCCTCCGGTA CCTACGCTAG CGACGCCGGC TAGAATCGGT CTGCTCGCCC
-----
4851 TTCGGCCCAT TCGGACCGCA AGGAATCGGT CAATACACTA CATGGCGTGA
      AAGCCGGGTA AGCCTGGCGT TCCTTAGCCA GTTATGTGAT GTACCGCACT
-----
4901 TTTCATATGC GCGATTGCTG ATCCCCATGT GTATCACTGG CAAACTGTGA
      AAAGTATACG CGCTAACGAC TAGGGGTACA CATAGTGACC GTTTGACACT
-----
4951 TGGACGACAC CGTCAGTGCG TCCGTCGCGC AGGCTCTCGA TGAGCTGATG
      ACCTGCTGTG GCAGTCACGC AGGCAGCGCG TCCGAGAGCT ACTCGACTAC
-----
5001 CTTTGGGCGG AGGACTGCCC CGAAGTCCGG CACCTCGTGC ACGCGGATTT
      GAAACCCGGC TCCTGACGGG GCTTCAGGCC GTGGAGCACG TCGCCTAAA
-----
5051 CGGCTCCAAC AATGTCCTGA CGGACAATGG CCGCATAACA GCGGTCATTG
      GCCGAGGTTG TTACAGGACT GCCTGTTACC GCGGTATTGT CGCCAGTAAC
-----
5101 ACTGGAGCGA GGCGATGTTT GGGGATTCCC AATACGAGGT CGCCAACATC
      TGACCTCGCT CCGCTACAAG CCCCTAAGGG TTATGCTCCA GCGGTTGTAG
-----
5151 TTCTTCTGGA GGCCGTGGTT GGCTTGTATG GAGCAGCAGA CGCGCTACTT
      AAGAAGACCT CCGGCACCAA CCGAACATAC CTCGTGCTCT GCGCGATGAA
-----
5201 CGAGCGGAGG CATCCGGAGC TTGCAGGATC GCCGCGGCTC CGGGCGTATA
      GCTCGCTCC GTAGGCCTCG AACGTCCTAG CGGCGCCGAG GCCCGCATAT
-----
5251 TGCTCCGCAT TGGTCTTGAC CAACTCTATC AGAGCTTGGT TGACGGCAAT
      ACGAGGCGTA ACCAGAACTG GTTGAGATAG TCTCGAACCA ACTGCCGTTA
-----
5301 TTCGATGATG CAGCTTGGGC GCAGGGTCGA TGCGACGCAA TCGTCCGATC
      AAGCTACTAC GTCGAACCCG CGTCCCAGCT ACGCTGCGTT AGCAGGCTAG
-----
5351 CGGAGCCGGG ACTGTCGGGC GTACACAAAT CGCCCGCAGA AGCGCGGCCG
      GCCTCGGCCC TGACAGCCCG CATGTGTTTA GCGGGCGTCT TCGCGCCGGC
-----
5401 TCTGGACCGA TGGCTGTGTA GAAGTACTCG CCGATAGTGG AAACCGACGC
      AGACCTGGCT ACCGACACAT CTTCATGAGC GGCTATCACC TTTGGCTGCG
-----
5451 CCCAGCACTC GTCCGAGGGC AAAGGAATAG AGTAGATGCC GACCGGGATC
      GGGTCGTGAG CAGGCTCCCG TTTCCTTATC TCATCTACGG CTGGCCCTAG
-----
5501 TATCGATAAA ATAAAAGATT TTATTTAGTC TCCAGAAAAA GGGGGGAATG
      ATAGCTATTT TATTTTCTAA AATAAATCAG AGGTCTTTTT CCCCCCTTAC
-----
5551 AAAGACCCCA CCTGTAGGTT TGGCAAGCTA GCTTAAGTAA CGCCATTTTG
      TTTCTGGGGT GGACATCCAA ACCGTTTCGAT CGAATTCATT GCGGTAAAC
-----
5601 CAAGGCATGG AAAAATACAT AACTGAGAAT AGAGAAGTTC AGATCAAGGT
      GTTCCGTACC TTTTATGTA TTGACTCTTA TCTCTTCAAG TCTAGTTCCA
-----
5651 CAGGAACAGA TGGAACAGCT GAATATGGGC CAAACAGGAT ATCTGTGGTA
      GTCCTTGTCT ACCTTGTCGA CTTATACCCG GTTTGTCCTA TAGACACCAT
-----

```



```

5701 AGCAGTTCCT GCGGCGGCTC AGGGCCAAGA ACAGATGGAA CAGCTGAATA
TCGTCAAGGA CGGGGCCGAG TCCCAGTTCT TGTCTACCTT GTCGACTTAT
-----
5751 TGGGCCAAAC AGGATATCTG TGGTAAGCAG TTCCTGCCCC GGCTCAGGGC
ACCCGGTTTG TCCTATAGAC ACCATTTCGT CAGGACGGGG CCGAGTCCCG
-----
5801 CAAGAACAGA TGGTCCCCAG ATGCGGTCCA GCCCTCAGCA GTTCTAGAG
GTTCTTGTCT ACCAGGGGTC TACGCCAGGT CGGGAGTCGT CAAAGATCTC
-----
5851 AACCATCAGA TGTTCCTCAG GTGCCCAAG GACCTGAAAT GACCCTGTGC
TTGGTAGTCT ACAAAGGTCC CACGGGGTTC CTGGACTTTA CTGGGACACG
-----
5901 CTTATTTGAA CTAACCAATC AGTTCGCTTC TCGCTTCTGT TCGCGCGCTT
GAATAAACTT GATTGGTTAG TCAAGCGAAG AGCGAAGACA AGCGCGCGAA
-----
5951 CTGCTCCCCG AGCTCAATAA AAGAGCCCAC AACCCCTCAC TCGGGGCGCC
GACGAGGGGC TCGAGTTATT TTCTCGGGTG TTGGGGAGTG AGCCCCGCGG
-----
6001 AGTCCTCCGA TTGACTGAGT CGCCCGGGTA CCCGTGTATC CAATAAACCC
TCAGGAGGCT AACTGACTCA GCGGGCCCAT GGGCACATAG GTTATTTGGG
-----
6051 TCTTGCAAGT GCATCCGACT TGTGGTCTCG CTGTTCCTTG GGAGGTCTC
AGAACGTCAA CGTAGGCTGA ACACCAGAGC GACAAGGAAC CCTCCCAGAG
-----
6101 CTCTGAGTGA TTGACTACCC GTCAGCGGGG GTCTTTCATT CATGCAGCAT
GAGACTCACT AACTGATGGG CAGTCGCCCC CAGAAAGTAA GTACGTCGTA
-----
6151 GTATCAAAAT TAATTTGGTT TTTTCTTTA AGTATTTACA TTAAATGGCC
CATAGTTTAA ATTAACCAA AAAAAAGAAT TCATAAATGT AATTACCGG
-----
6201 ATAGTTGCAT TAATGAATCG GCCAACGCGC GGGGAGAGGC GGTTCGCGTA
TATCAACGTA ATTACTTAGC CGGTTGCGCG CCCCTCTCCG CCAAACGCAT
-----
6251 TTGGCGCTCT TCCGCTTCCT CGCTCACTGA CTCGCTGCGC TCGGTCGTTC
AACCGCGAGA AGGCGAAGGA GCGAGTGACT GAGCGACGCG AGCCAGCAAG
-----
6301 GGCTGCGGCG AGCGGTATCA GCTCACTCAA AGGCGGTAAT ACGGTTATCC
CCGACGCCGC TCGCCATAGT CGAGTGAGTT TCCGCCATTA TGCCAATAGG
-----
6351 ACAGAATCAG GGGATAACGC AGGAAAGAAC ATGTGAGCAA AAGGCCAGCA
TGTCTTAGTC CCCTATTGCG TCCTTTCTTG TACACTCGTT TTCCGTCGT
-----
6401 AAAGGCCAGG AACCGTAAAA AGGCCGCGTT GCTGGCGTTT TTCCATAGGC
TTTCCGGTCC TTGGCATTTC TCCGGCGCAA CGACCGCAA AAGGTATCCG
-----
6451 TCCGCCCCCC TGACGAGCAT CACAAAAATC GACGCTCAAG TCAGAGGTGG
AGGCGGGGGG ACTGCTCGTA GTGTTTTTAG CTGCGAGTTC AGTCTCCACC
-----
6501 CGAAACCCGA CAGGACTATA AAGATACCAG GCGTTTCCCC CTGGAAGCTC
GCTTTGGGCT GTCCTGATAT TTCTATGGTC CGCAAAGGGG GACCTTCGAG
-----
6551 CCTCGTGCGC TCTCCTGTTT CGACCCTGCC GCTTACCGGA TACCTGTCCG
GGAGCACGCG AGAGGACAAG GCTGGGACGG CGAATGGCCT ATGGACAGGC
-----
6601 CCTTTCTCCC TTCGGGAAGC GTGGCGCTTT CTCATAGCTC ACGCTGTAGG
GGAAAGAGGG AAGCCCTTCG CACCGCGAAA GAGTATCGAG TGCGACATCC
-----

```

```
6651 TATCTCAGTT CGGTGTAGGT CGTTCGCTCC AAGCTGGGCT GTGTGCACGA
      ATAGAGTCAA GCCACATCCA GCAAGCGAGG TTCGACCCGA CACACGTGCT
-----
6701 ACCCCCCGTT CAGCCCCACC GCTGCGCCTT ATCCGGTAAC TATCGTCTTG
      TGGGGGGCAA GTCGGGCTGG CGACGCGGAA TAGGCCATTG ATAGCAGAAC
-----
6751 AGTCCAACCC GGTAAGACAC GACTTATCGC CACTGGCAGC AGCCACTGGT
      TCAGGTTGGG CCATTCTGTG CTGAATAGCG GTGACCGTCG TCGGTGACCA
-----
6801 AACAGGATTA GCAGAGCGAG GTATGTAGGC GGTGCTACAG AGTTCTTGAA
      TTGTCTAAT CGTCTCGCTC CATACATCCG CCACGATGTC TCAAGAACTT
-----
6851 GTGGTGGCCT AACTACGGCT AACTAGAAAG AACAGTATTT GGTATCTGCG
      CACCACCGGA TTGATGCCGA TGTGATCTTC TTGTCATAAA CCATAGACGC
-----
6901 CTCTGCTGAA GCCAGTTACC TTCGGAAAAA GAGTTGGTAG CTCTTGATCC
      GAGACGACTT CGGTCAATGG AAGCCTTTTT CTCAACCATC GAGAACTAGG
-----
6951 GGCAAACAAA CCACCGCTGG TAGCGGTGGT TTTTGTGTTT GCAAGCAGCA
      CCGTTTGTGT GGTGGCGACC ATCGCCACCA AAAAAACAAA CGTTCGTCGT
-----
7001 GATTACGCGC AGAAAAAAG GATCTCAAGA AGATCCTTTG ATCTTTTCTA
      CTAATGCGCG TCTTTTTTTC CTAGAGTTCT TCTAGGAAAC TAGAAAAGAT
-----
7051 CGGGGTCTGA CGCTCAGTGG AACGAAACT CACGTTAAGG GATTTTGGTC
      GCCCAGACT GCGAGTCACC TTGCTTTTGA GTGCAATTCC CTAAAACCAG
-----
7101 ATGAGATTAT CAAAAAGGAT CTTACCTAG ATCCTTTTGC GGCCGCAAAT
      TACTCTAATA GTTTTCTCTA GAAGTGGATC TAGGAAACG CCGGCGTTTA
-----
7151 CAATCTAAAG TATATATGAG TAACTTGGT CTGACAGTGA CCAATGCTTA
      GTTAGATTTC ATATATACTC ATTTGAACCA GACTGTCAAT GGTTACGAAT
-----
7201 ATCAGTGAGG CACCTATCTC AGCGATCTGT CTATTTCTGT CATCCATAGT
      TAGTCACTCC GTGGATAGAG TCGCTAGACA GATAAAGCAA GTAGGTATCA
-----
7251 TGCCTGACTC CCCGTCGTGT AGATAACTAC GATACGGGAG GGCTTACCAT
      ACGGACTGAG GGGCAGCACA TCTATTGATG CTATGCCCTC CCGAATGGTA
-----
7301 CTGGCCCCAG TGCTGCAATG ATACCGCGAG ACCCAGCTC ACCGGCTCCA
      GACCGGGGTC ACGACGTTAC TATGGCGCTC TGGGTGCGAG TGGCCGAGGT
-----
7351 GATTTATCAG CAATAAACCA GCCAGCCGGA AGGGCCGAGC GCAGAAGTGG
      CTAAATAGTC GTTATTTGGT CGGTCGGCCT TCCCGGCTCG CGTCTTCACC
-----
7401 TCCTGCAACT TTATCCGCTT CCATCCAGTC TATTAATTGT TGCCGGGAAG
      AGGACGTTGA AATAGGCGGA GGTAGGTCAG ATAATTAACA ACGGCCCTTC
-----
7451 CTAGAGTAAG TAGTTCGCCA GTTAATAGTT TGCGCAACGT GTTGCCATT
      GATCTCATTC ATCAAGCGGT CAATTATCAA ACGCGTTGCA ACAACGGTAA
-----
7501 GCTACAGGCA TCGTGGTGTC ACGCTCGTCG TTTGGTATGG CTTCAATCAG
      CGATGTCCGT AGCACCACAG TGCGAGCAGC AAACCATACC GAAGTAAGTC
-----
7551 CTCCGTTTCC CAACGATCAA GGCGAGTTAC ATGATCCCCC ATGTTGTGCA
      GAGGCCAAGG GTTGCTAGTT CCGCTCAATG TACTAGGGGG TACAACACGT
-----
```

7601 AAAAAGCGGT TAGCTCCTTC GGTCTCCGA TCGTTGTCAG AAGTAAGTTG  
TTTTTCGCCA ATCGAGGAAG CCAGGAGGCT AGCAACAGTC TTCATTCAAC

7651 GCCGCAGTGT TATCACTCAT GGTATGGCA GCACTGCATA ATTCTCTTAC  
CGGCGTCACA ATAGTGAGTA CCAATACCGT CGTGACGTAT TAAGAGAATG

7701 TGTCATGCCA TCCGTAAGAT GCTTTTCTGT GACTGGTGAG TACTCAACCA  
ACAGTACGGT AGGCATTCTA CGAAAAGACA CTGACCACTC ATGAGTTGGT

7751 AGTCATTCTG AGAATAGTGT ATGCGGCGAC CGAGTTGCTC TTGCCCCGCG  
TCAGTAAGAC TCTTATCACA TACGCCGCTG GCTCAACGAG AACGGGCCGC

7801 TCAATACGGG ATAATACCGC GCCACATAGC AGAACTTTAA AAGTGCTCAT  
AGTTATGCCC TATTATGGCG CCGGTGTATCG TCTTGAAATT TTCACGAGTA

7851 CATTGGAAAA CGTCTTTCGG GCGGAAACT CTCAAGGATC TTACCGCTGT  
GTAACCTTTT GCAAGAAGCC CCGCTTTTGA GAGTTCCTAG AATGGCGACA

7901 TGAGATCCAG TTCGATGTAA CCCACTCGTG CACCCAACCTG ATCTTCAGCA  
ACTCTAGGTC AAGCTACATT GGGTGAGCAC GTGGGTTGAC TAGAAGTCGT

7951 TCTTTTACTT TCACCAGCGT TTCTGGGTGA GCAAAAACAG GAAGGCAAAA  
AGAAAATGAA AGTGGTCGCA AAGACCCACT CGTTTTTGTC CTTCCGTTTT

8001 TGCCGCAAAA AAGGGAATAA GGGCGACACG GAAATGTTGA ATACTCATAC  
ACGGCGTTTT TTCCCTTATT CCCGCTGTGC CTTTACAACT TATGAGTATG

8051 TCTTCCTTTT TCAATATTAT TGAAGCATTT ATCAGGGTTA TTGTCTCATG  
AGAAGGAAAA AGTTATAATA ACTTCGTAAA TAGTCCAAT AACAGAGTAC

8101 AGCGGATACA TATTTGAATG TATTTAGAAA AATAAACAAA TAGGGGTTCC  
TCGCCTATGT ATAAACTTAC ATAAATCTTT TTATTTGTTT ATCCCCAAGG

8151 GCGCACATTT C  
CGCGTGATAA G

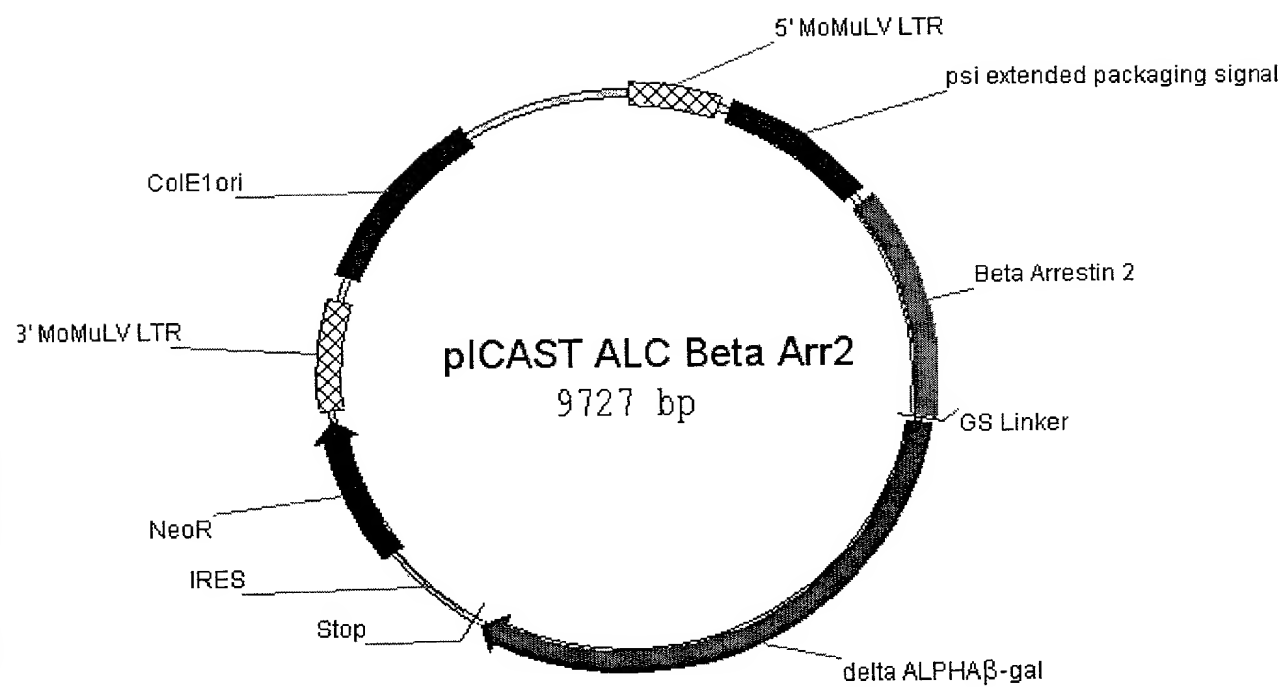


Figure 14

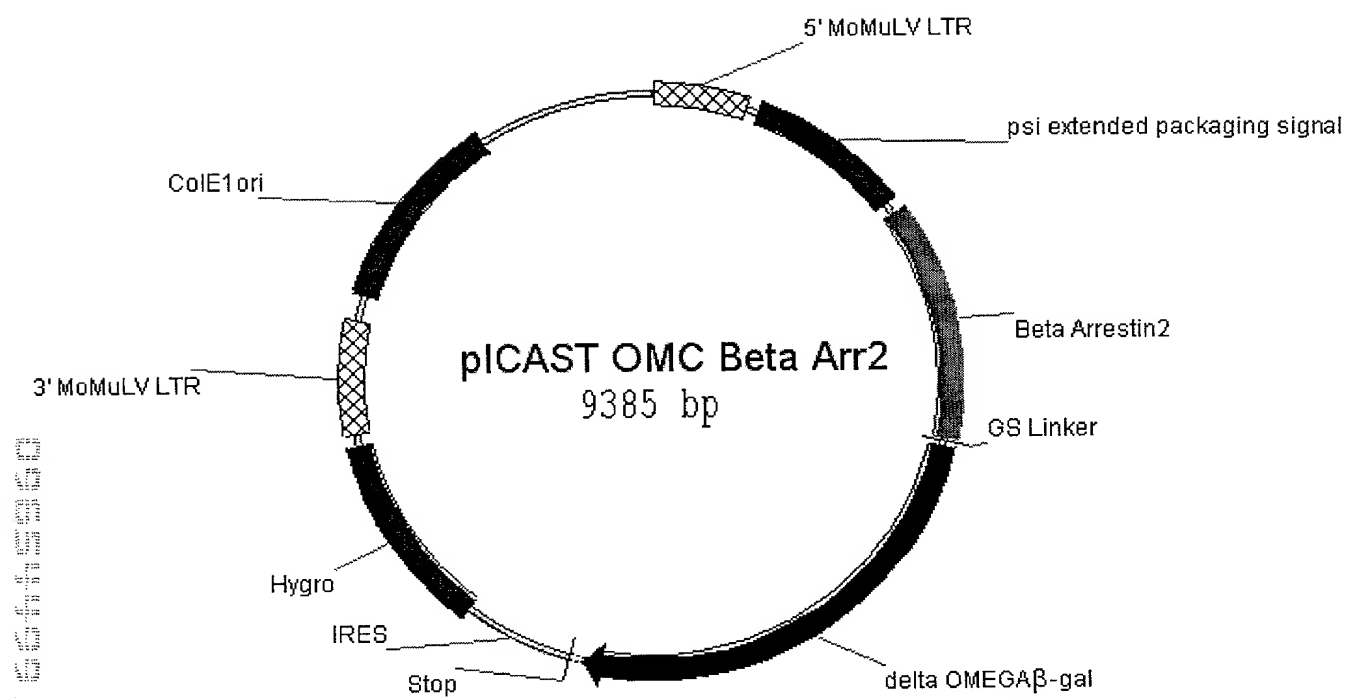


Figure 15

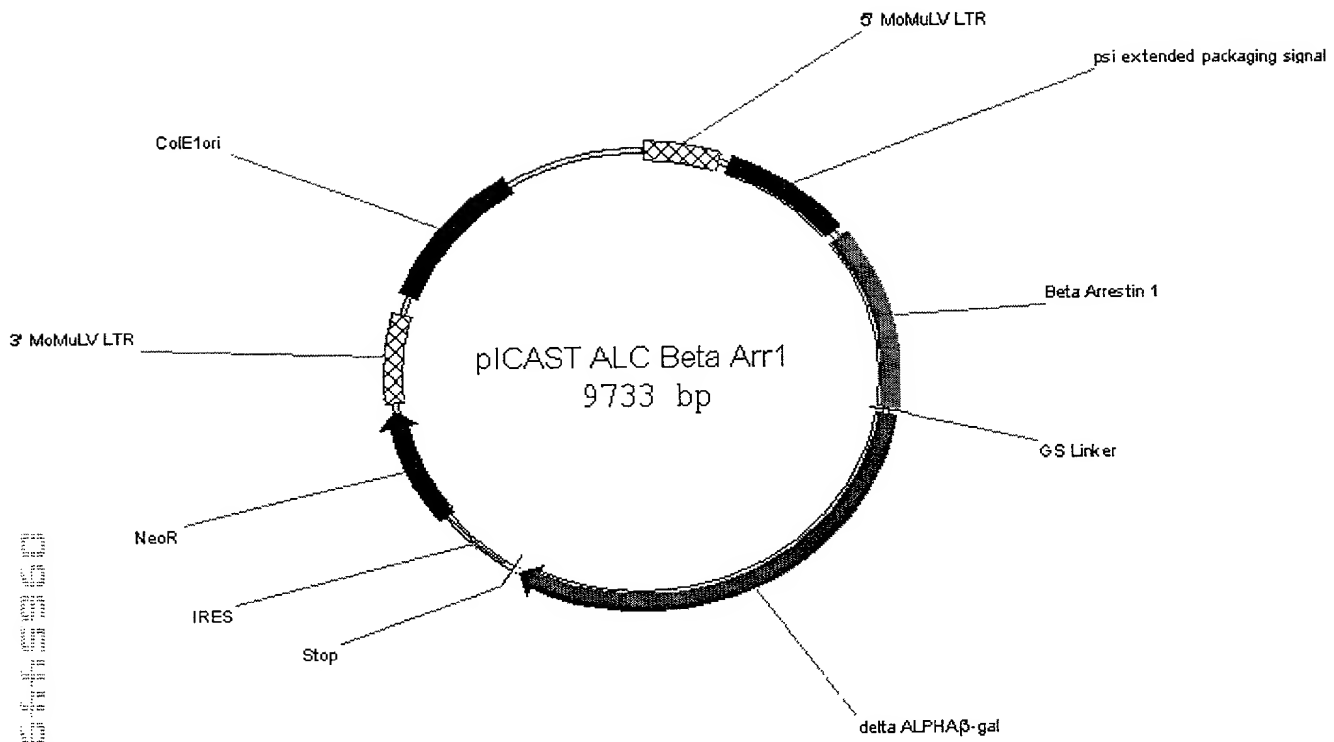


Figure 16

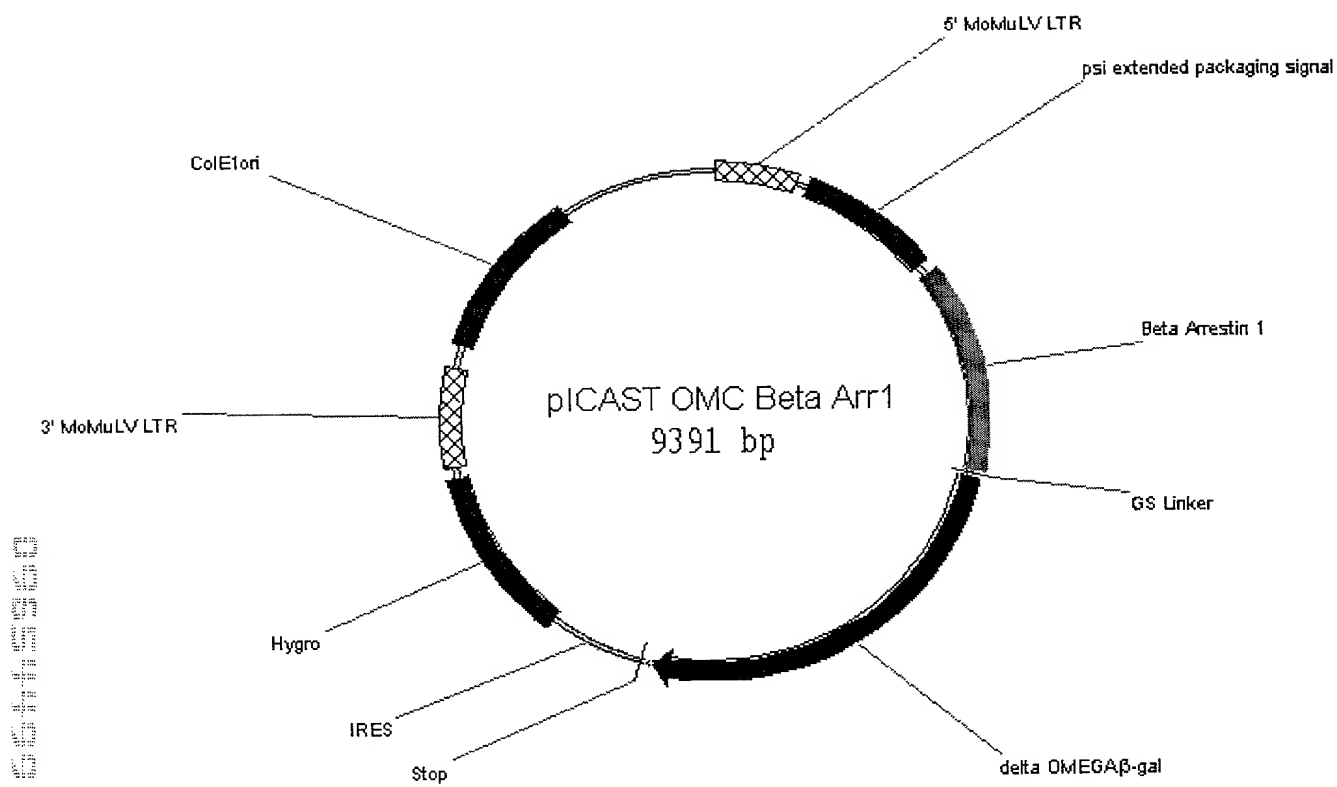


Figure 17

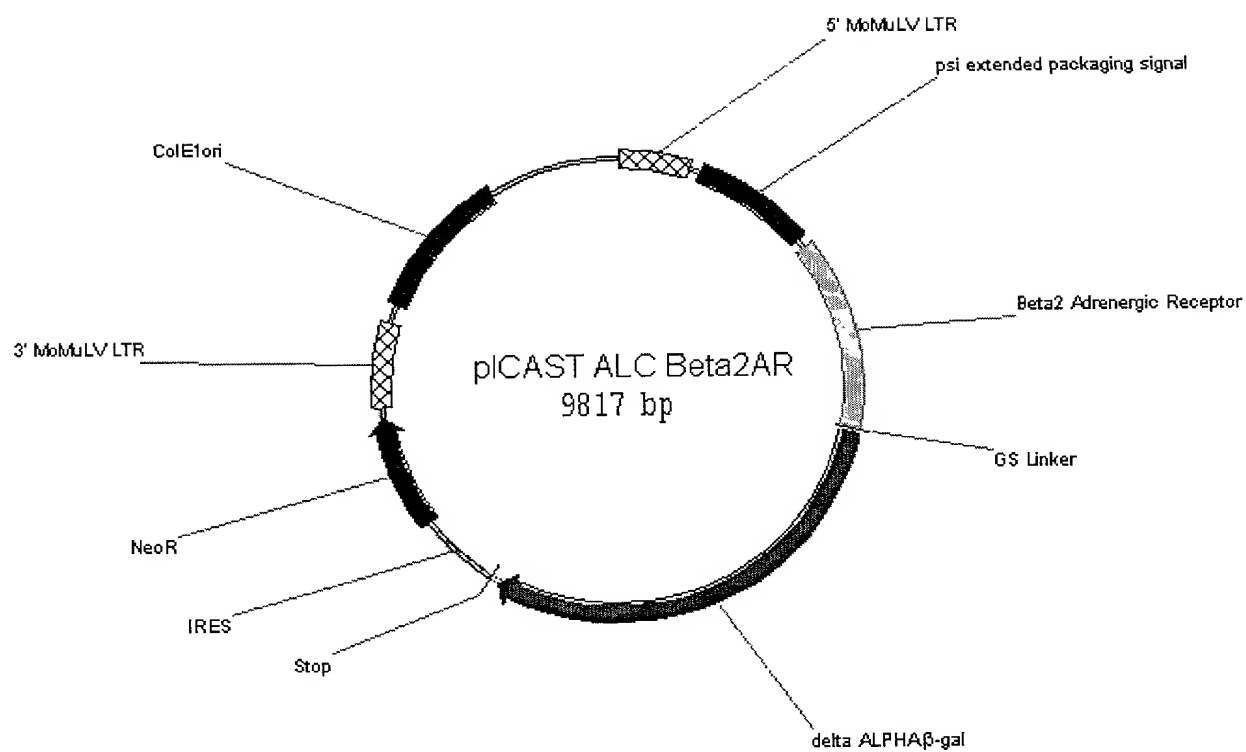


Figure 18



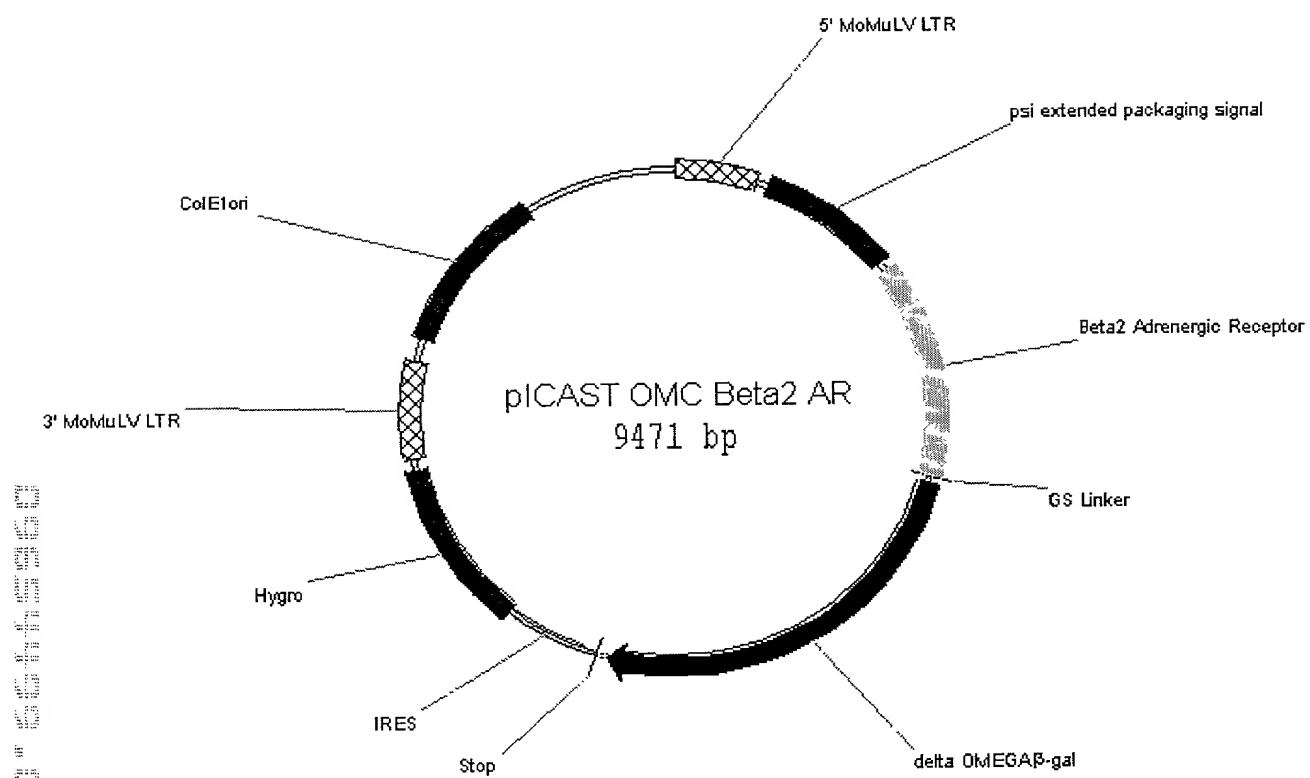


Figure 19

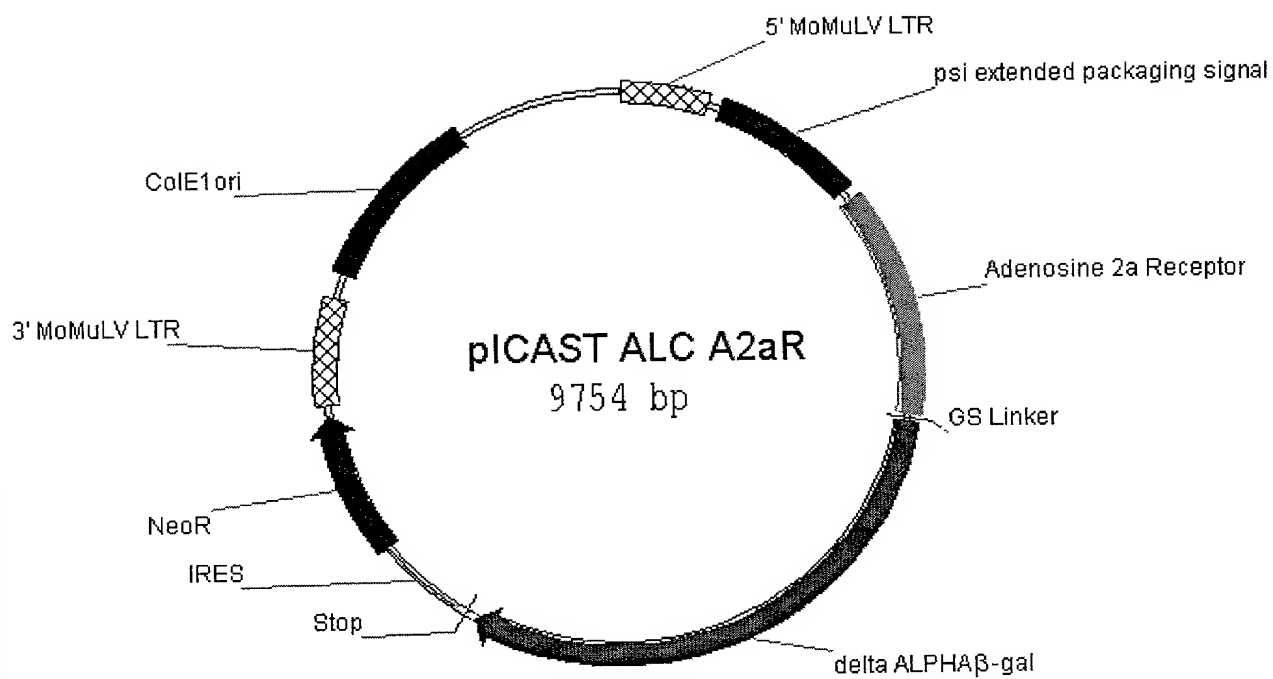


Figure 20

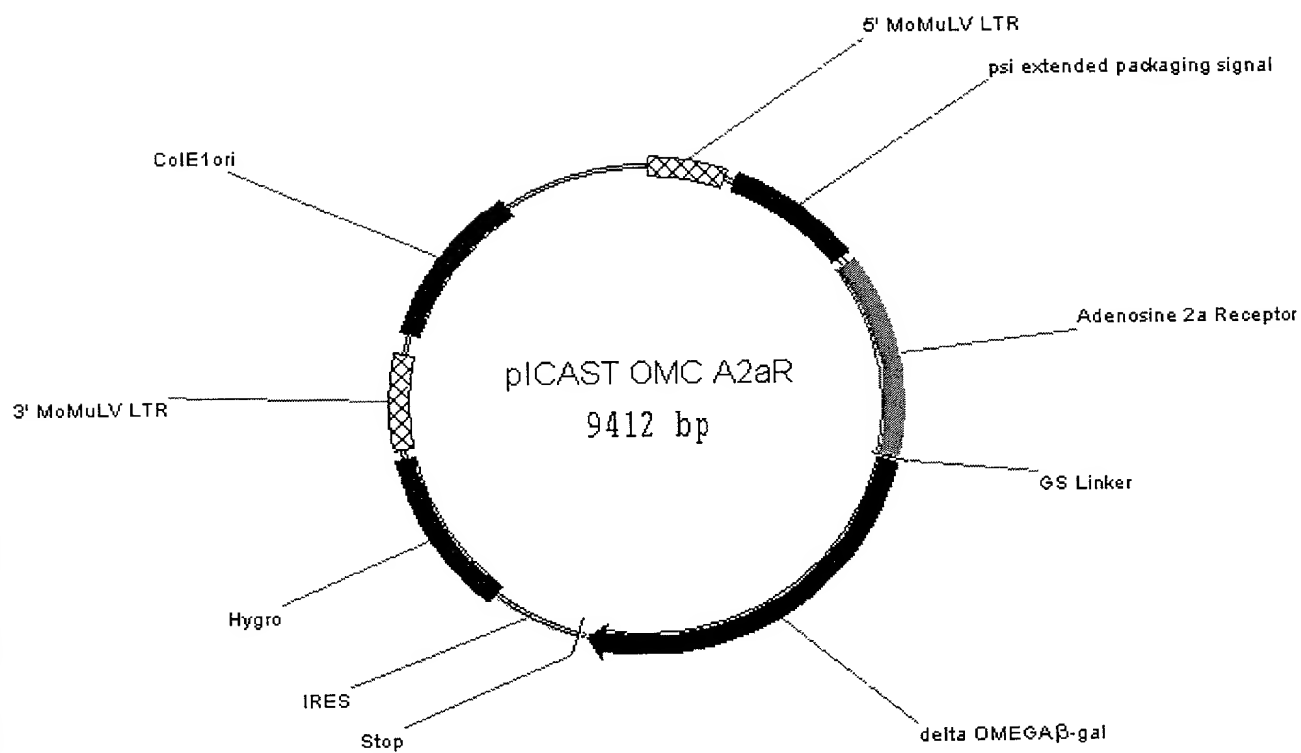


Figure 21

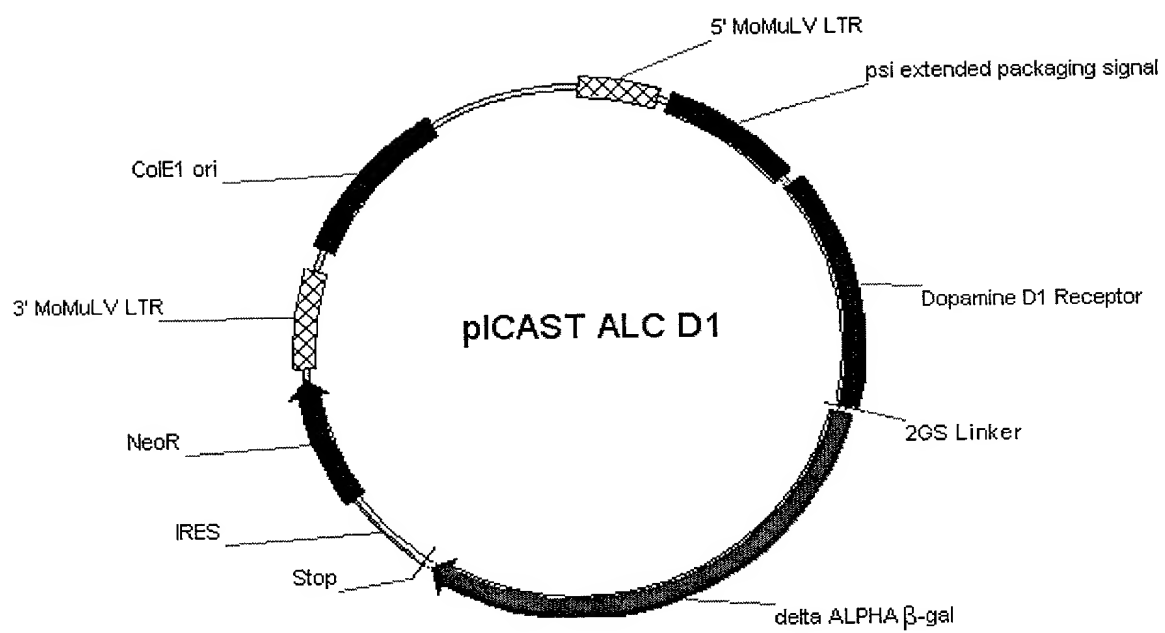
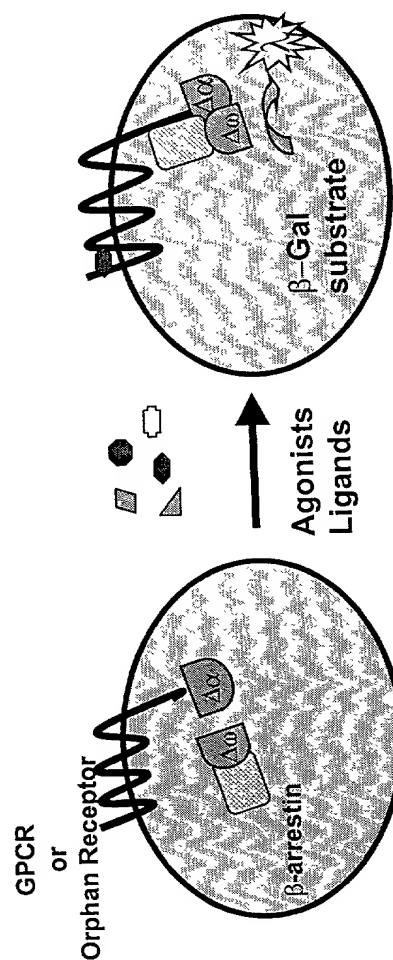


Figure 22

Functional GPCR Activation Assay and Ligand Fishing for Orphan Receptors  
by  $\beta$ -galactosidase mutant complementation in ICAS<sup>TM</sup> System



Examples

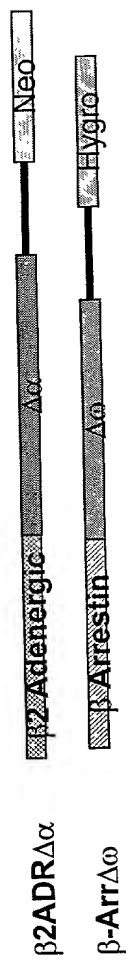


Figure 23

DOCKET NO. 4085-226-27

**IN THE UNITED STATES PATENT AND TRADEMARK OFFICE**

IN RE APPLICATION OF: Michelle A.J. PALMER, et al

ART UNIT:

SERIAL NO.: New Application

EXAMINER:

FILING DATE: Herewith

FOR: RECEPTOR FUNCTION ASSAY FOR G-PROTEIN COUPLED  
RECEPTORS AND ORPHAN RECEPTORS BY REPORTER ENZYME  
MUTANT COMPLEMENTATION

**LIST OF INVENTORS' NAMES AND ADDRESSES**

ASSISTANT COMMISSIONER FOR PATENTS  
WASHINGTON, D.C. 20231

SIR:


Listed below are the names and addresses of the inventors for the above-identified patent application.

- |   |  |
|---|--|
| 1) Michelle A.J. PALMER<br>87 Medford Street<br>Arlington, MA 02174 | 3) Bonnie TILLOTSON<br>4 Ripley Road<br>Belmont, MA 02178    |
| 2) Melissa GEE<br>3 North Street<br>Grafton, MA 01519               | 4) Xiao-Jia CHANG<br>25 Round Hill Road<br>Lincoln, MA 01773 |

A declaration containing all the necessary information will be submitted at a later date.

Respectfully submitted,

PIPER MARBURY RUDNICK & WOLFE LLP

  
\_\_\_\_\_  
Steven B. Kelber  
Registration No. 30,073  
Attorney of Record

Laura A. Donnelly  
Registration No. 38,435

1200 Nineteenth Street, N.W.  
Washington, D.C. 20036-2412  
Telephone No. (202) 861-3900  
Facsimile No. (202) 223-2085